

PCT

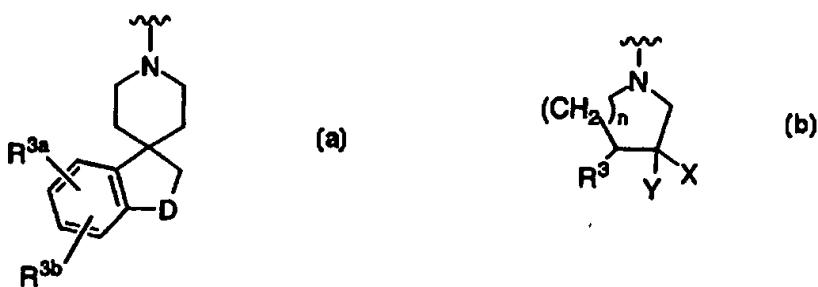
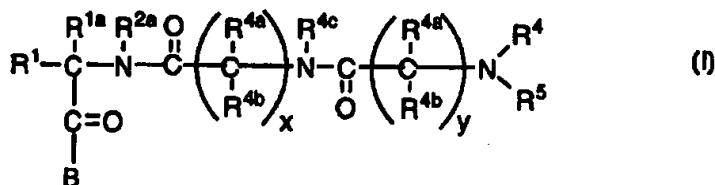
WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : C07D 221/20, 401/12, 401/14, A61K 31/40, 31/445		A1	(11) International Publication Number: WO 97/36873			
			(43) International Publication Date: 9 October 1997 (09.10.97)			
(21) International Application Number:	PCT/US97/05378		(74) Common Representative: MERCK & CO., INC.; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).			
(22) International Filing Date:	31 March 1997 (31.03.97)					
(30) Priority Data:			(81) Designated States: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, HU, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, US, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).			
(71) Applicant (for all designated States except US):	MERCK & CO., INC. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).					
(72) Inventors; and						
(75) Inventors/Applicants (for US only):	CHAKRAVARTY, Prasun, K. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). CHEN, Meng, H. [Stateless/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). NARGUND, Ravi [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). PATCHETT, Arthur, A. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). TATA, James, R. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). WU, Mu, Tsu [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). YANG, Lihu [CN/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).					
Published						
With international search report.						
Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.						

(54) Title: PIPERIDINES, PYRROLIDINES AND HEXAHYDRO-1H-AZEPINES PROMOTE RELEASE OF GROWTH HORMONE



(57) Abstract

The present invention is directed to certain piperidines, pyrrolidines, and hexahydro-1H-azepines of general structural formula (I), wherein B is selected from (a) and (b), and R¹, R^{1a}, R^{2a}, R^{3a}, R^{3b}, R⁴, R^{4a}, R^{4b}, R^{4c}, R⁵, D, X, Y, n, x and y are as defined herein. These compounds promote the release of growth hormone in humans and animals. This property can be utilized to promote the growth of food animals to render the production of edible meat products more efficient, and in humans, to treat physiological or medical conditions characterized by a deficiency in growth hormone secretion, such as short stature in growth hormone deficient children, and to treat medical conditions which are improved by the anabolic effects of growth hormone. Growth hormone releasing compositions containing such compounds as the active ingredient thereof are also disclosed.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LJ	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

TITLE OF THE INVENTION

PIPERIDINES, PYRROLIDINES AND HEXAHYDRO-1H-AZEPINES PROMOTE RELEASE OF GROWTH HORMONE

5 BACKGROUND OF THE INVENTION

Growth hormone, which is secreted from the pituitary, stimulates growth of all tissues of the body that are capable of growing. In addition, growth hormone is known to have the following basic effects on the metabolic processes of the body: (1) Increased rate of protein synthesis in all cells of the body; (2) Decreased rate of carbohydrate utilization in cells of the body; (3) Increased mobilization of free fatty acids and use of fatty acids for energy. A deficiency in growth hormone secretion can result in various medical disorders, such as dwarfism.

15 Various ways are known to release growth hormone. For example, chemicals such as arginine, L-3,4-dihydroxyphenylalanine (L-DOPA), glucagon, vasopressin, and insulin induced hypoglycemia, as well as activities such as sleep and exercise, indirectly cause growth hormone to be released from the pituitary by

20 acting in some fashion on the hypothalamus perhaps either to decrease somatostatin secretion or to increase the secretion of the known secretagogue growth hormone releasing factor (GRF) or an unknown endogenous growth hormone-releasing hormone or all of these.

25 In cases where increased levels of growth hormone were desired, the problem was generally solved by providing exogenous growth hormone or by administering GRF or a peptidal compound which stimulated growth hormone production and/or release. In either case the peptidyl nature of the compound necessitated that it be

30 administered by injection. Initially the source of growth hormone was the extraction of the pituitary glands of cadavers. This resulted in a very expensive product and carried with it the risk that a disease associated with the source of the pituitary gland could be transmitted to the recipient of the growth hormone. Recombinant growth

- 2 -

hormone has become available which, while no longer carrying any risk of disease transmission, is still a very expensive product which must be given by injection or by a nasal spray. Other compounds have been developed which stimulate the release of endogenous 5 growth hormone such as analogous peptidyl compounds related to GRF or the peptides of U.S. Patent 4,411,890. These peptides, while considerably smaller than growth hormones are still susceptible to various proteases. As with most peptides, their potential for oral bioavailability is low. Non peptidal growth hormone secretagogues 10 are disclosed in e.g., U.S. Patent Nos 5,206,235, 5,283,241, 5,284,841, 5,310,737, 5,317,017, 5,374,721, 5,430,144, 5,434,261, 5,438,136, 5,494,919, 5,494,920, and 5,492,916. Other growth 15 hormone secretagogues are disclosed e.g., in PCT Patent Publications WO 94/13696, WO 94/19367, WO 95/03289, WO 95/03290, WO 95/09633, WO 95/11029, WO 95/12598, WO 95/13069, WO 95/14666, WO 95/16675, WO 95/16692, WO 95/17422, WO 95/17423, WO 95/34311, and WO 96/02530. The instant compounds 20 are low molecular weight peptide analogs for promoting the release of growth hormone which have good stability in a variety of physiological environments and which may be administered parenterally, nasally or by the oral route.

SUMMARY OF THE INVENTION

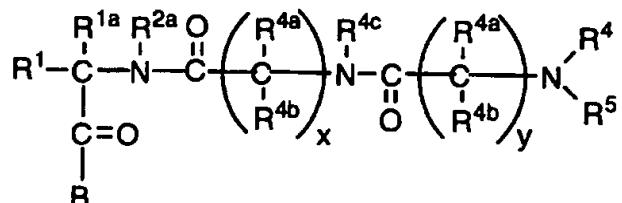
The instant invention is directed to certain piperidines, 25 pyrrolidines, and hexahydro-1H-azepines which have the ability to stimulate the release of natural or endogenous growth hormone. The compounds thus have the ability to be used to treat conditions which require the stimulation of growth hormone production or secretion such as in humans with a deficiency of natural growth hormone or in animals 30 used for food or wool production where the stimulation of growth hormone will result in a larger, more productive animal. Thus, it is an object of the instant invention to describe the piperidine compounds. It is a further object of this invention to describe procedures for the preparation of such compounds. A still further object is to describe the

- 3 -

use of such compounds to increase the secretion of growth hormone in humans and animals. A still further object of this invention is to describe compositions containing the piperidine compounds for the use of treating humans and animals so as to increase the level of growth hormone secretions. Further objects will become apparent from a reading of the following description.

DESCRIPTION OF THE INVENTION

10 The novel piperidines, pyrrolidines, and hexahydro-1H-azepines of the instant invention are described by structural Formula I:



Formula I

15 wherein:

R¹ is selected from the group consisting of:
C₁-C₁₀ alkyl, -aryl-, aryl (C₁-C₆ alkyl)-,
heteroaryl-, heteroaryl(C₁-C₆ alkyl)-,
(C₃-C₇ cycloalkyl)-(C₁-C₆ alkyl)-,
20 (C₁-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
aryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
heteroaryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-, and
(C₃-C₇ cycloalkyl)-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
wherein K is -O-, -S(O)_m-, -N(R²)C(O)-, -C(O)N(R²)-, -OC(O)-,
25 -C(O)O-, -CR²=CR²- or -C≡C-,
wherein R² and the alkyl groups may be further substituted with 1 to 9
halo, -S(O)_mR^{2a}, 1 to 3 of -OR^{2a}, or -C(O)OR^{2a}, and wherein aryl is
phenyl or naphthyl, and heteroaryl is selected from indolyl, thiophenyl,
benzofuranyl, benzothiophenyl, aza-indolyl, pyridinyl, quinolinyl, and
30 benzimidazolyl, wherein aryl and heteroaryl are unsubstituted or
substituted with phenyl, phenoxy, halophenyl, 1 to 3 of -C₁-C₆ alkyl, 1

- 4 -

to 3 of halo, 1 to 2 of -OR², methylenedioxy, -S(O)_mR², 1 to 2 of -CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²), -C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, or -N(R²)SO₂R²;

5 R^{1a} is hydrogen or C₁-C₄ alkyl;

R² is selected from the group consisting of:

hydrogen, -C₁-C₆ alkyl, -C₃-C₇ cycloalkyl, and -CH₂-phenyl,

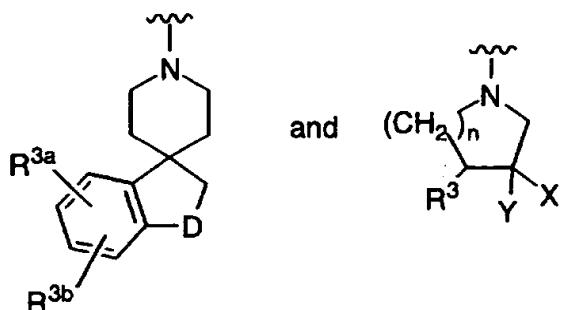
wherein the alkyl or the cycloalkyl is unsubstituted or substituted with

10 hydroxyl, C₁-C₃ alkoxy, thioalkyl, C(O)OR^{2a}, and where, if two -C₁-C₆ alkyl groups are present on one atom, they may be joined to form a C₃-C₈ cyclic ring being selected from the group consisting of pyrrolidine, piperidine, piperazine, morpholine, thiomorpholine, optionally substituted by hydroxyl;

15

R^{2a} is hydrogen or C₁-C₆ alkyl;

B is selected from:



20 R³ is selected from: hydrogen, -(CH₂)_rphenyl, -(CH₂)_rthienyl, -(CH₂)_rbenzimidazolyl, -(CH₂)_rquinolinyl, -(CH₂)_rnaphthyl, -(CH₂)_rindolyl, -C₁-C₁₀ alkyl, -C₃-C₇ cycloalkyl, where the phenyl, naphthyl, indolyl, thienyl, benzimidazolyl, quinolinyl, and C₃-C₇ cycloalkyl rings may be substituted by 1 to 3 substituents selected from

25 the group consisting of: C₁-C₆ alkyl, halogen, -OR², -NHSO₂CF₃, -(CH₂)_rOR⁶, -(CH₂)_rN(R²)(R⁶), -(CH₂)_r(R⁶), -(CH₂)_rC(O)OR², -(CH₂)_rC(O)OR⁶, -(CH₂)_rOC(O)R², -(CH₂)_rOC(O)R⁶,

- 5 -

- $(CH_2)_rC(O)R^2$, $-(CH_2)_rC(O)R^6$, $-(CH_2)_rC(O)N(R^2)(R^2)$,
- $(CH_2)_rC(O)N(R^2)(R^6)$, $-(CH_2)_rN(R^2)C(O)(R^2)$,
- $(CH_2)_rN(R^2)C(O)R^6$ $-(CH_2)_rN(R^6)C(O)R^2$, $-(CH_2)_rN(R^6)C(O)R^6$,
- $(CH_2)_rN(R^2)C(O)OR^2$, $-(CH_2)_rN(R^2)C(O)OR^6$,
- 5 $-(CH_2)_rN(R^6)C(O)OR^2$, $-(CH_2)_rN(R^6)C(O)OR^6$,
- $(CH_2)_rN(R^2)C(O)N(R^2)(R^6)$, $-(CH_2)_rN(R^2)C(O)N(R^2)(R^2)$,
- $(CH_2)_rN(R^6)C(O)N(R^2)(R^6)$, $-(CH_2)_rN(R^2)SO_2R^2$,
- $(CH_2)_rN(R^6)SO_2R^2$, $-(CH_2)_rN(R^6)SO_2R^6$, $-(CH_2)_rOC(O)N(R^2)(R^6)$,
- $(CH_2)_rOC(O)N(R^2)(R^2)$, $-(CH_2)_rSO_2N(R^2)(R^6)$,
- 10 $-(CH_2)_rOC(O)N(R^2)(R^2)$, $-(CH_2)_rSO_2N(R^2)(R^6)$,
- $(CH_2)_rSO_2N(R^2)(R^2)$, $-(CH_2)_rN(R^2)SO_2N(R^2)(R^6)$,
- $(CH_2)_rN(R^6)SO_2N(R^2)(R^6)$, $-(CH_2)_rS(O)_mR^6$, and $-(CH_2)_rS(O)_mR^2$;

- R^{3a} and R^{3b} are independently selected from: hydrogen, phenyl, phenoxy, halophenyl, -C₁-C₆ alkyl, halogen, -OR², methylenedioxy, -S(O)_mR², -CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²), -C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, and -N(R²)SO₂R²;
- 20 R⁴ and R⁵ are independently selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from halo, hydroxy, phenyl, and C₁-C₆ alkoxy carbonyl; or R⁵ and R⁴ may be taken together to form -(CH₂)_d-La(CH₂)_e- where La is -C(R²)₂-, -O-, -S(O)_m- or -N(R²)-, d and e are independently 1 to 3 and R² is as defined above;

- R^{4a} and R^{4b} are independently selected from: hydrogen, C₁-C₆ alkyl, trifluoromethyl, phenyl, or substituted C₁-C₆ alkyl where the substituents are selected from: imidazolyl, naphthyl, phenyl, indolyl, p-hydroxyphenyl, -OR², -S(O)_mR², -C(O)OR², C₃-C₇ cycloalkyl, -N(R²)(R²), -C(O)N(R²)(R²); or R^{4a} and R^{4b} may independently be joined to one or both of R⁴ or R^{4c} to form an alkylene bridge between the terminal nitrogen and the alkyl portion of the R^{4a} or R^{4b} and the

- 6 -

R^4 or R^{4c} group, wherein the bridges contain 1 to 8 carbons atoms; or R^{4a} and R^{4b} may be joined to one another to form C3-C7 cycloalkyl;

R^{4c} is selected from hydrogen, C1-C6 alkyl, and substituted C1-C6

5 alkyl where the substituents are selected from 1 to 5 halo, 1 to 3 hydroxy, phenyl, naphthyl, imidazolyl, indolyl, and C1-C6 alkoxy carbonyl;

10 R^6 is selected from: hydrogen, C1-C6 alkyl, and $(CH_2)_q$ aryl, wherein the $(CH_2)_q$ and alkyl groups may be optionally substituted by $-O(R^2)$, $-S(O)_mR^2$, $-C(O)OR^2$, $-C(O)N(R^2)(R^2)$, $-SO_2N(R^2)(R^2)$, or $-N(R^2)C(O)N(R^2)(R^2)$, wherein the aryl group is selected from: phenyl, pyridyl, 1H-tetrazolyl, triazolyl, oxadiazolyl, pyrazolyl, thiadiazolyl, and benzimidazol-2-yl, which is optionally substituted with 15 C1-C6 alkyl, C3-C6 cycloalkyl, amino, or hydroxyl;

20 X is selected from the group consisting of: hydrogen, $-C\equiv N$, $-(CH_2)_qN(R^2)C(O)R^2$, $-(CH_2)_qN(R^2)C(O)(CH_2)_t$ aryl, $-(CH_2)_qN(R^2)SO_2(CH_2)_t$ aryl, $-(CH_2)_qN(R^2)SO_2R^2$, $-(CH_2)_qN(R^2)C(O)N(R^2)(CH_2)_t$ aryl, $-(CH_2)_qN(R^2)C(O)N(R^2)(R^2)$, $-(CH_2)_qC(O)N(R^2)(R^2)$, $-(CH_2)_qC(O)N(R^2)(CH_2)_t$ aryl, $-(CH_2)_qC(O)OR^2$, $-(CH_2)_qC(O)O(CH_2)_t$ aryl, $-(CH_2)_qOR^2$, $-(CH_2)_qOC(O)R^2$, $-(CH_2)_qOC(O)(CH_2)_t$ aryl, $-(CH_2)_qOC(O)N(R^2)(R^2)$, $-(CH_2)_qC(O)R^2$, $-(CH_2)_qC(O)(CH_2)_t$ aryl, 25 $-(CH_2)_qN(R^2)C(O)OR^2$, $-(CH_2)_qN(R^2)SO_2N(R^2)(R^2)$, $-(CH_2)_qS(O)_mR^2$, and $-(CH_2)_qS(O)_m(CH_2)_t$ aryl, where R^2 , $(CH_2)_q$ and $(CH_2)_t$ group may be optionally substituted with C1-C4 alkyl, hydroxyl, C1-C4 lower alkoxy, carboxyl, $N(R^2)(R^2)$, $CONH_2$, $S(O)_mCH_3$, carboxylate C1-C4 alkyl esters, or 1H-tetrazol-5-yl, and 30 aryl is phenyl, naphthyl, pyridyl, thiazolyl, or 1H-tetrazol-5-yl groups which may be optionally substituted with halogen, $-OR^2$, $-CON(R^2)(R^2)$, $-C(O)OR^2$, C1-C4 alkyl, $-S(O)_mR^2$, or 1H-tetrazol-5-yl;

- 7 -

Y is selected from the group consisting of:

hydrogen, C₁-C₁₀ alkyl, -(CH₂)_taryl,
 -(CH₂)_q(C₃-C₇ cycloalkyl), -(CH₂)_q-K-(C₁-C₆ alkyl), -(CH₂)_q-K-(CH₂)_taryl, -(CH₂)_q-K-(CH₂)_t(C₃-C₇ cycloalkyl containing O, NR² S)

- 5 and -(CH₂)_q-K-(CH₂)_t(C₃-C₇ cycloalkyl), where K is O, S(O)_m,
 C(O)NR², CH=CH, C≡C, N(R²)C(O), C(O)NR², C(O)O, or OC(O),
 and where the alkyl, R², (CH₂)_q and (CH₂)_t groups are optionally
 substituted by C₁-C₄ alkyl, hydroxyl, C₁-C₄ lower alkoxy, carboxyl,
 -CONH₂ or a carboxylate C₁-C₄ alkyl ester, and aryl is phenyl,
- 10 naphthyl, pyridyl, 1-H-tetrazol-5-yl, thiazolyl, imidazolyl, indolyl,
 oxadiazolyl, pyrimidinyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl,
 thiophenyl, quinolinyl, pyrazinyl, or isothiazolyl which is optionally
 substituted with halogen, -OR², -C(O)OR², N(R²)(R²),
 -C(O)N(R²)(R²), nitro, cyano, benzyl,
- 15 C₁-C₄ alkyl, -S(O)_mR², or 1H-tetrazol-5-yl;

D is selected from: -N(R⁷)-, -S(O)_m-, -C(O)- and -C(H)(R⁷)-,
 wherein R⁷ is selected from: -R², -OR², -(CH₂)_qaryl, -C(O)R²,
 -C(O)(CH₂)_qaryl, -SO₂R², -SO₂(CH₂)_qaryl, -C(O)N(R²)(R²),

- 20 -C(O)N(R²)(CH₂)_qaryl, -C(O)OR², 1-H-tetrazol-5-yl, -SO₂N(R²)aryl,
 -SO₂N(R²)(R²) and the (CH₂)_q may be optionally substituted by C₁-C₄
 alkyl, and the R² and aryl may be optionally further substituted with a
 substituent selected from: -OR^{2a}, -O(CH₂)_q aryl, -C(O)OR^{2a},
 -C(O)(CH₂)_q aryl, -C(O)N(R^{2a})(R^{2a}), -C(O)N(R^{2a})(CH₂)₁ aryl,
- 25 halogen, -N(R^{2a})(R^{2a}), -C₁-C₄ alkyl, 1,2,4-triazolyl, 1-H-tetrazol-5-yl,
 -C(O)NHSO₂R^{2a}, -S(O)_mR^{2a}, -C(O)NHSO₂(CH₂)_qaryl,
 -N(R²)C(O)N(R^{2a})(R^{2a}), -N(R^{2a})C(O)N(R^{2a})(CH₂)_qaryl,
 -N(R^{2a})(R^{2a}), -N(R^{2a})C(O)R^{2a}, -N(R^{2a})C(O)(CH₂)_q aryl,
 -OC(O)N(R^{2a})(R^{2a}), -OC(O)N(R^{2a})(CH₂)_q aryl;

- 30
 - 1 is 0, 1 or 2;
 - m is 0, 1, or 2;
 - n is 0, 1, or 2;
 - q is 0, 1, 2, 3, or 4;

- 8 -

r is 0, 1, 2, or 3;
t is 0, 1, 2, or 3;
v is 0, 1, or 2;

5 and pharmaceutically acceptable salts and individual diastereomers thereof.

In the above structural formula and throughout the instant specification, the following terms have the indicated meanings:

10

When n is 1 a pyrrolidine ring is formed, when n is 2 a piperidine ring is formed, and when n is 3 the ring is designated a hexahydro-1-H-azepine.

15

The alkyl groups specified above are intended to include those alkyl groups of the designated length in either a straight or branched configuration and if two carbon atoms or more they may include a double or a triple bond. Exemplary of such alkyl groups are methyl, ethyl, propyl, isopropyl, butyl, sec-butyl, tertiary butyl, pentyl, isopentyl, hexyl, isohexyl, allyl, propargyl, and the like.

20

The alkoxy groups specified above are intended to include those alkoxy groups of the designated length in either a straight or branched configuration and if two or more carbon atoms in length, they may include a double or a triple bond. Exemplary of such alkoxy groups are methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, tertiary butoxy, pentoxy, isopentoxy, hexoxy, isohexoxy allyloxy, propargyloxy, and the like.

25

The term "halogen" is intended to include the halogen atom fluorine, chlorine, bromine and iodine.

30

The term "aryl" within the present invention, unless otherwise specified, is intended to include aromatic rings, such as carbocyclic and heterocyclic aromatic rings selected the group consisting of: phenyl, naphthyl, pyridyl, 1-H-tetrazol-5-yl, thiazolyl, imidazolyl, indolyl, pyrimidinyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl, thiophenyl, quinolinyl, pyrazinyl, or isothiazolyl, which

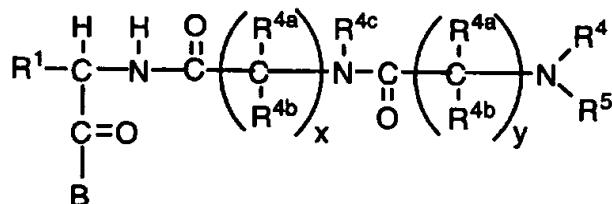
- 9 -

may be optionally substituted by 1 to 3 of C₁-C₆ alkyl, 1 to 3 of halogen, 1 to 2 of OR₂, methylenedioxy, -S(O)_mR₂, 1 to 2 of -CF₃, -OCF₃, nitro, -N(R₂)C(O)(R₂), -C(O)OR₂, -C(O)N(R₂)(R₂), -1H-tetrazol-5-yl, -SO₂N(R₂)(R₂), -N(R₂)SO₂ phenyl, or -N(R₂)SO₂R₂,

5 wherein R₂ is as defined herein.

Certain of the above defined terms may occur more than once in the above formula and upon such occurrence each term shall be defined independently of the other.

10 Preferred compounds of the instant invention include those of Formula Ia:



Formula Ia

wherein:

15 R¹ is selected from the group consisting of:
 C₁-C₁₀ alkyl, -aryl-, aryl (C₁-C₆ alkyl)-,
 heteroaryl-, heteroaryl(C₁-C₆ alkyl)-,
 (C₃-C₇ cycloalkyl)-(C₁-C₆ alkyl)-,
 (C₁-C₅ alkyl)-K-(C₁-C₅ alkyl)-,

20 aryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 heteroaryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-, and
 (C₃-C₇ cycloalkyl)-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 wherein K is -O-, -S(O)_m-, -N(R²)C(O)-, -C(O)N(R²)-, -OC(O)-,
 -C(O)O-, -CR²=CR²- or -C≡C-,

25 wherein R² and the alkyl groups may be further substituted with 1 to 9 halo, -S(O)_mR^{2a}, 1 to 3 of -OR^{2a}, or -C(O)OR^{2a}, and wherein aryl is phenyl or naphthyl, and heteroaryl is selected from indolyl, thiophenyl, benzofuranyl, benzothiophenyl, aza-indolyl, pyridinyl, quinolinyl, and benzimidazolyl, wherein aryl and heteroaryl are unsubstituted or

30 substituted with phenyl, phenoxy, halophenyl, 1 to 3 of -C₁-C₆ alkyl, 1 to 3 of halo, 1 to 2 of -OR², methylenedioxy, -S(O)_mR², 1 to 2 of

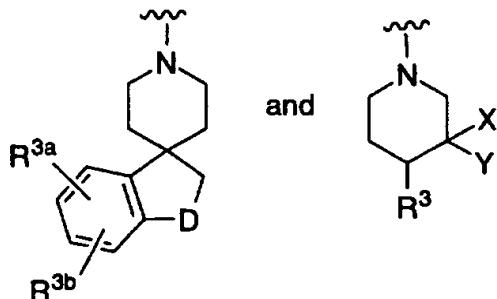
- 10 -

-CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²), -C(O)OR²,
-C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, or -N(R²)SO₂R²;

R² is selected from the group consisting of:

5 hydrogen, -C₁-C₆ alkyl, -C₃-C₇ cycloalkyl, and -CH₂-phenyl,
wherein the alkyl or the cycloalkyl is unsubstituted or substituted with
hydroxyl, C₁-C₃ alkoxy, thioalkyl, -C(O)OR^{2a}, and wherein, if two
-C₁-C₆ alkyl groups are present on one atom, the groups may be
optionally joined to form a C₃-C₈ cyclic ring being selected from the
10 group consisting of pyrrolidine, piperidine, piperazine, morpholine,
thiomorpholine;

B is selected from:



15 R³ is selected from: hydrogen, phenyl, naphthyl, indolyl,
benzimidazolyl, thienyl, quinolinyl, where the phenyl, naphthyl,
benzimidazolyl, thienyl, quinolinyl, and indolyl may be substituted by 1
to 3 substituents selected from the group consisting of: C₁-C₆ alkyl,
halogen, -OR², -(CH₂)_rOR⁶, -(CH₂)_rN(R²)(R⁶), -(CH₂)_r(R⁶),
-(CH₂)_rC(O)OR², -(CH₂)_rC(O)OR⁶, -(CH₂)_rC(O)R², -(CH₂)_rC(O)R⁶,
20 -(CH₂)_rC(O)N(R²)(R²), -(CH₂)_rC(O)N(R²)(R⁶),
-(CH₂)_rN(R²)C(O)(R²), -(CH₂)_rN(R²)C(O)R⁶, -(CH₂)_rN(R⁶)C(O)R²,
-(CH₂)_rN(R⁶)C(O)R⁶, -(CH₂)_rN(R²)C(O)OR²,
-(CH₂)_rN(R²)C(O)OR⁶, -(CH₂)_rN(R⁶)C(O)OR²,
25 -(CH₂)_rN(R⁶)C(O)OR⁶, -(CH₂)_rN(R²)C(O)N(R²)(R⁶),
-(CH₂)_rN(R²)C(O)N(R²)(R²), -(CH₂)_rN(R⁶)C(O)N(R²)(R⁶),
-(CH₂)_rN(R²)SO₂R², -(CH₂)_rN(R⁶)SO₂R², -(CH₂)_rN(R⁶)SO₂R⁶,

- 11 -

$-(CH_2)_rOC(O)N(R^2)(R^6)$, $-(CH_2)_rSO_2N(R^2)(R^6)$,
 $-(CH_2)_rSO_2N(R^2)(R^6)$, $-(CH_2)_rSO_2N(R^2)(R^2)$, $-(CH_2)_rS(O)_mR^6$, and
 $-(CH_2)_rS(O)_mR^2$;

5 R^{3a} and R^{3b} are independently selected from: hydrogen, phenyl, phenoxy, halophenyl, -C₁-C₆ alkyl, halogen, -OR², methylenedioxy, -S(O)_mR², -CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²), -C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, and -N(R²)SO²R²;

10 10 R^4 and R^5 are independently selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from halo, hydroxy, phenyl, and C₁-C₆ alkoxy carbonyl; or R^5 and R^4 may be taken together to form -(CH₂)_d-La(CH₂)_e- where

15 15 La is -C(R²)₂- , -O- , -S(O)_m- or -N(R²)-, d and e are independently 1 to 3 and R² is as defined above;

20 R^{4a} and R^{4b} are independently selected from: hydrogen, C₁-C₆ alkyl, trifluoromethyl, phenyl, or substituted C₁-C₆ alkyl where the substituents are selected from: imidazolyl, naphthyl, phenyl, indolyl, p-hydroxyphenyl, -OR², -S(O)_mR², -C(O)OR², C₃-C₇ cycloalkyl, -N(R²)(R²), -C(O)N(R²)(R²); or R^{4a} and R^{4b} may independently be joined to one or both of R^4 or R^{4c} to form an alkylene bridge between the terminal nitrogen and the alkyl portion of the R^{4a} or R^{4b} and the

25 25 R^4 or R^{4c} group, wherein the bridges contain 1 to 5 carbon atoms; or R^{4a} and R^{4b} may be joined to one another to form C₃-C₇ cycloalkyl;

30 R^{4c} is selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from 1 to 5 halo, 1 to 3 hydroxy, phenyl, C₁-C₆ alkoxy carbonyl;

35 R^6 is selected from: hydrogen, C₁-C₆ alkyl, and (CH₂)_varyl, wherein the (CH₂)_v and alkyl groups may be optionally substituted by -O(R²), -S(O)_mR², -C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), or

- 12 -

$-\text{N}(\text{R}^2)\text{C}(\text{O})\text{N}(\text{R}^2)(\text{R}^2)$, wherein the aryl group is selected from: phenyl, pyridyl, 1H-tetrazolyl, triazolyl, oxadiazolyl, pyrazolyl, thiadiazoyl, and benzimidazol-2-yl, which is optionally substituted with C1-C6 alkyl, C3-C6 cycloalkyl, amino, or hydroxyl;

5

X is selected from the group consisting of: hydrogen, $-\text{C}\equiv\text{N}$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{C}(\text{O})\text{R}^2$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{C}(\text{O})(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{SO}_2(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{SO}_2\text{R}^2$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{C}(\text{O})\text{N}(\text{R}^2)(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{C}(\text{O})\text{N}(\text{R}^2)(\text{R}^2)$,

10 $-(\text{CH}_2)_q\text{C}(\text{O})\text{N}(\text{R}^2)(\text{R}^2)$, $-(\text{CH}_2)_q\text{C}(\text{O})\text{N}(\text{R}^2)(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{C}(\text{O})\text{OR}^2$, $-(\text{CH}_2)_q\text{C}(\text{O})\text{O}(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{OR}^2$, $-(\text{CH}_2)_q\text{OC}(\text{O})\text{R}^2$, $-(\text{CH}_2)_q\text{OC}(\text{O})(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{OC}(\text{O})\text{N}(\text{R}^2)(\text{R}^2)$, $-(\text{CH}_2)_q\text{C}(\text{O})\text{R}^2$, $-(\text{CH}_2)_q\text{C}(\text{O})(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{C}(\text{O})\text{OR}^2$, $-(\text{CH}_2)_q\text{N}(\text{R}^2)\text{SO}_2\text{N}(\text{R}^2)(\text{R}^2)$,

15 $-(\text{CH}_2)_q\text{S}(\text{O})_m\text{R}^2$, and $-(\text{CH}_2)_q\text{S}(\text{O})_m(\text{CH}_2)_t\text{aryl}$, where R^2 , $(\text{CH}_2)_q$ and $(\text{CH}_2)_t$ group may be optionally substituted with C1-C4 alkyl, hydroxyl, C1-C4 lower alkoxy, carboxyl, $\text{N}(\text{R}^2)(\text{R}^2)$, CONH_2 , $\text{S}(\text{O})_m\text{CH}_3$, carboxylate C1-C4 alkyl esters, or 1H-tetrazol-5-yl, and aryl is phenyl, naphthyl, pyridyl, thiazolyl, or 1H-tetrazol-5-yl groups

20 which may be optionally substituted with halogen, $-\text{OR}^2$, $-\text{CON}(\text{R}^2)(\text{R}^2)$, $-\text{C}(\text{O})\text{OR}^2$, C1-C4 alkyl, $-\text{S}(\text{O})_m\text{R}^2$, or 1H-tetrazol-5-yl;

Y is selected from the group consisting of:

25 hydrogen, C1-C10 alkyl, $-(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q(\text{C}_3\text{-C}_7\text{ cycloalkyl})$, $-(\text{CH}_2)_q\text{-K-}(\text{C}_1\text{-C}_6\text{ alkyl})$, $-(\text{CH}_2)_q\text{-K-}(\text{CH}_2)_t\text{aryl}$, $-(\text{CH}_2)_q\text{-K-}(\text{CH}_2)_t(\text{C}_3\text{-C}_7\text{ cycloalkyl containing O, NR}^2\text{ S})$ and $-(\text{CH}_2)_q\text{-K-}(\text{CH}_2)_t(\text{C}_3\text{-C}_7\text{ cycloalkyl})$, where K is O, $\text{S}(\text{O})_m$, $\text{C}(\text{O})\text{NR}^2$, $\text{CH}=\text{CH}$, $\text{C}\equiv\text{C}$, $\text{N}(\text{R}^2)\text{C}(\text{O})$, $\text{C}(\text{O})\text{NR}^2$, $\text{C}(\text{O})\text{O}$, or $\text{OC}(\text{O})$,

30 and where the alkyl, R^2 , $(\text{CH}_2)_q$ and $(\text{CH}_2)_t$ groups are optionally substituted by C1-C4 alkyl, hydroxyl, C1-C4 lower alkoxy, carboxyl, $-\text{CONH}_2$ or a carboxylate C1-C4 alkyl ester, and aryl is phenyl, naphthyl, pyridyl, 1-H-tetrazol-5-yl, thiazolyl, imidazolyl, indolyl, oxadiazoyl, pyrimidinyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl,

- 13 -

thiophenyl, quinolinyl, pyrrazinyl, or isothiazolyl which is optionally substituted with halogen, -OR², -C(O)OR², N(R²)(R²), -C(O)N(R²)(R²), nitro, cyano, benzyl, C₁-C₄ alkyl, -S(O)_mR², or 1H-tetrazol-5-yl;

5 D is selected from: -N(R⁷)-, -S(O)_m-, -C(O)- and -C(H)(R⁷)-, wherein R⁷ is selected from: -R², -(CH₂)_qaryl, -C(O)R², -SO₂R², -C(O)N(R²)(R²), -C(O)OR², 1-H-tetrazol-5-yl, -SO₂N(R²)aryl, -SO₂N(R²)(R²) and the (CH₂)_q may be optionally substituted by C₁-C₄ alkyl, and the R² and aryl may be optionally further substituted with a substituent selected from: -OR^{2a}, -C(O)OR^{2a}, -C(O)N(R^{2a})(R^{2a}), halogen, -C₁-C₄ alkyl, and the aryl is selected from of triazolyl, oxadiazolyl, thiadiazolyl, thiazolyl, imidazolyl, and 1H-tetrazolyl;

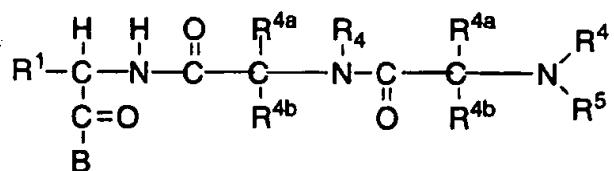
10 15 1 is 0, 1 or 2;
m is 0, 1, or 2;
q is 0, 1, 2, 3, or 4;
r is 0, 1, 2, or 3;
t is 0, 1, 2, or 3;

15 20 v is 0, 1, or 2;

and pharmaceutically acceptable salts and individual diastereomers thereof.

- 14 -

More preferred compounds of the instant invention include those of Formula Ib:

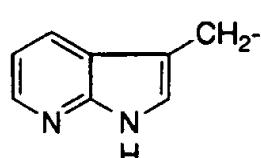
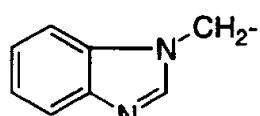
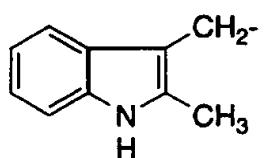
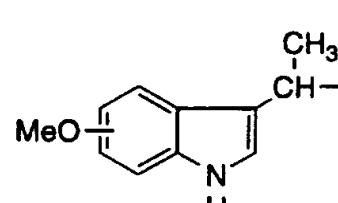
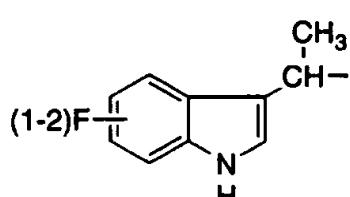
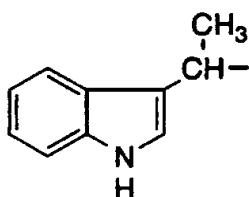
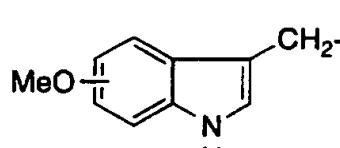
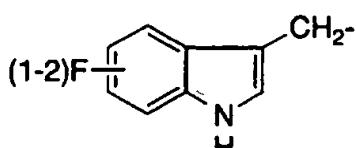
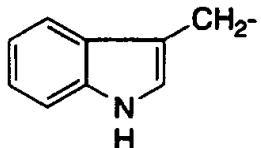


5

Formula Ib

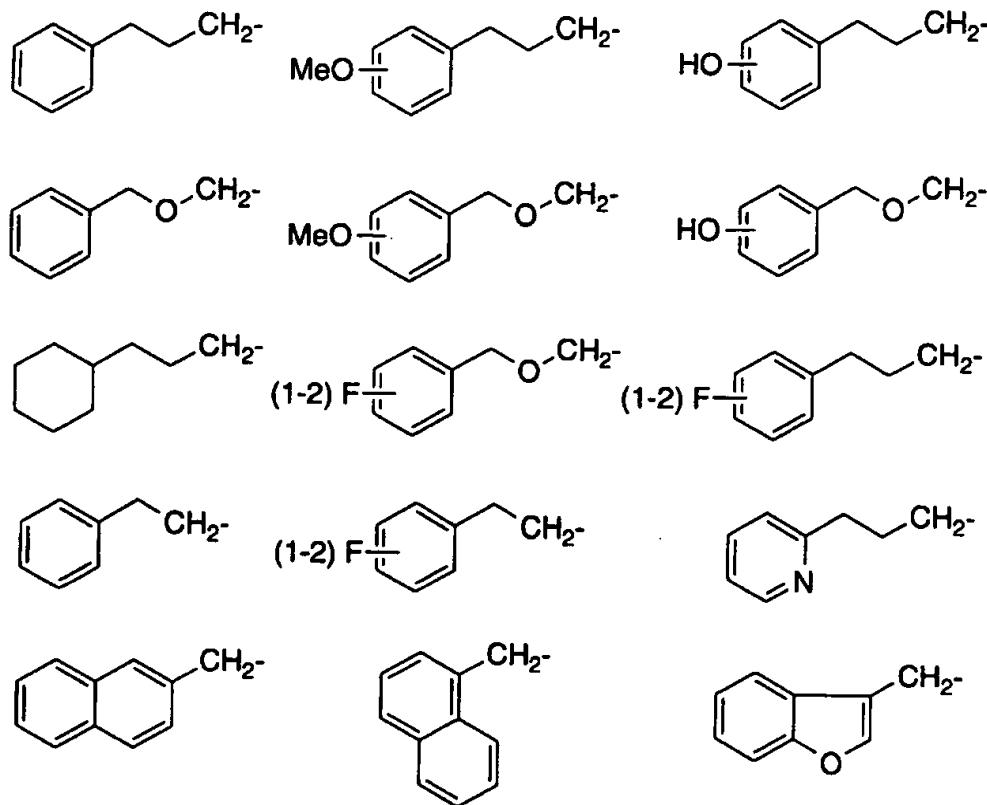
wherein:

R^1 is selected from the group consisting of:



10

- 15 -



or their regioisomers where not specified;

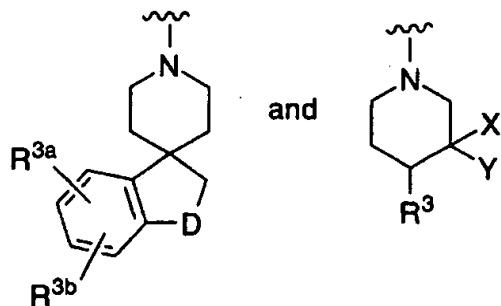
R² is selected from the group consisting of:

5 hydrogen, -C₁-C₆ alkyl, -C₃-C₇ cycloalkyl, and -CH₂-phenyl,
 wherein the alkyl or the cycloalkyl is unsubstituted or substituted with
 hydroxyl, C₁-C₃ alkoxy, thioalkyl, -C(O)OR^{2a}, and wherein, if two
 -C₁-C₆ alkyl groups are present on one atom, the groups may be
 optionally joined to form a C₃-C₈ cyclic ring being selected from the
 10 group consisting of pyrrolidine, piperidine, piperazine, morpholine,
 thiomorpholine;

R^{2a} is hydrogen, or C₁-C₄ alkyl;

- 16 -

B is selected from:



R³ is selected from: hydrogen or phenyl, wherein the phenyl is substituted in the ortho position by a substituent selected from the group consisting of: C₁-C₆ alkyl, halogen, -OR², -(CH₂)_rOR⁶, -(CH₂)_rN(R²)(R⁶), -(CH₂)_r(R⁶), -(CH₂)_rC(O)OR², -(CH₂)_rC(O)OR⁶, -(CH₂)_rC(O)R², -(CH₂)_rC(O)R⁶, -(CH₂)_rC(O)N(R²)(R²), -(CH₂)_rC(O)N(R²)(R⁶), -(CH₂)_rSO₂N(R²)(R⁶), -(CH₂)_rSO₂N(R²)(R²), -(CH₂)_rS(O)_mR⁶, and -(CH₂)_rS(O)_mR²;

10 **R**^{3a} and **R**^{3b} are independently selected from: hydrogen, -C₁-C₆ alkyl and halogen;

15 **R**⁴ and **R**⁵ are independently selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from halo, hydroxy, phenyl, and C₁-C₆ alkoxy carbonyl; or **R**⁵ and **R**⁴ may be taken together to form -(CH₂)_d-La(CH₂)_e- where La is -C(R²)₂- , -O-, -S(O)_m- or -N(R²)-, d and e are independently 1 to 3 and R² is as defined above;

20 **R**^{4a} and **R**^{4b} are independently selected from: hydrogen, C₁-C₆ alkyl, or substituted C₁-C₆ alkyl where the substituents are selected from: imidazolyl, naphthyl, phenyl, indolyl, and p-hydroxyphenyl;

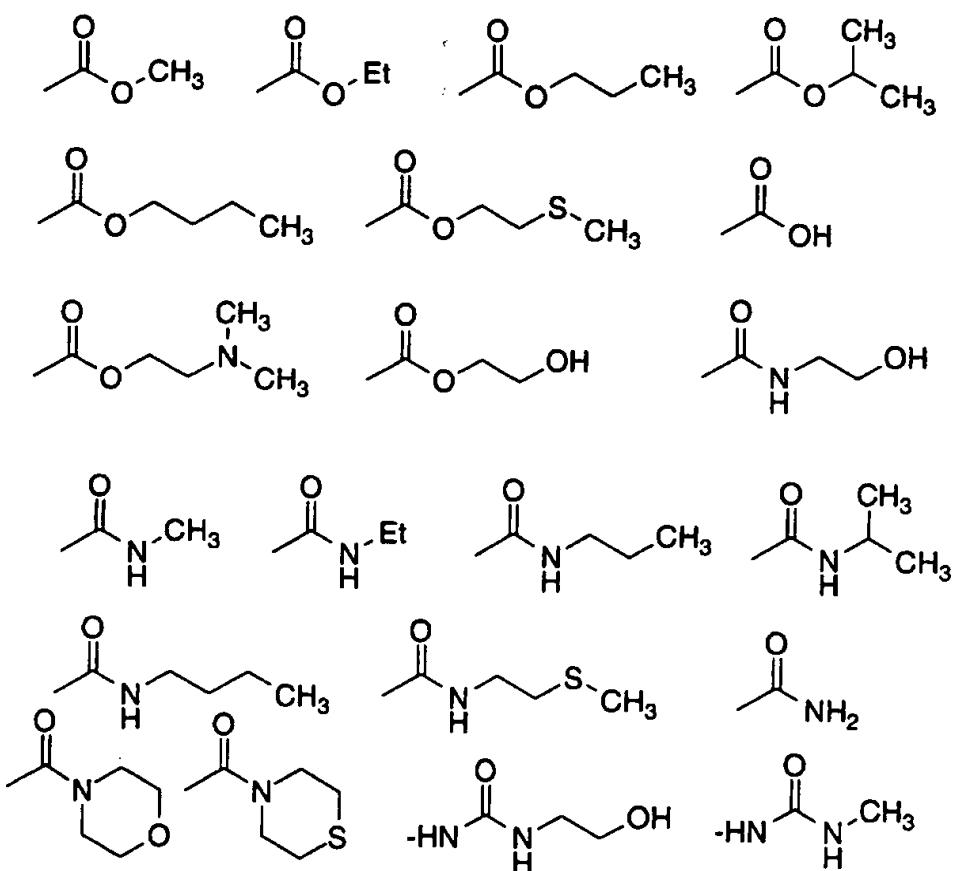
25 **R**^{4c} is selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from 1 to 5 halo, 1 to 3 hydroxy, phenyl, C₁-C₆ alkoxy carbonyl;

- 17 -

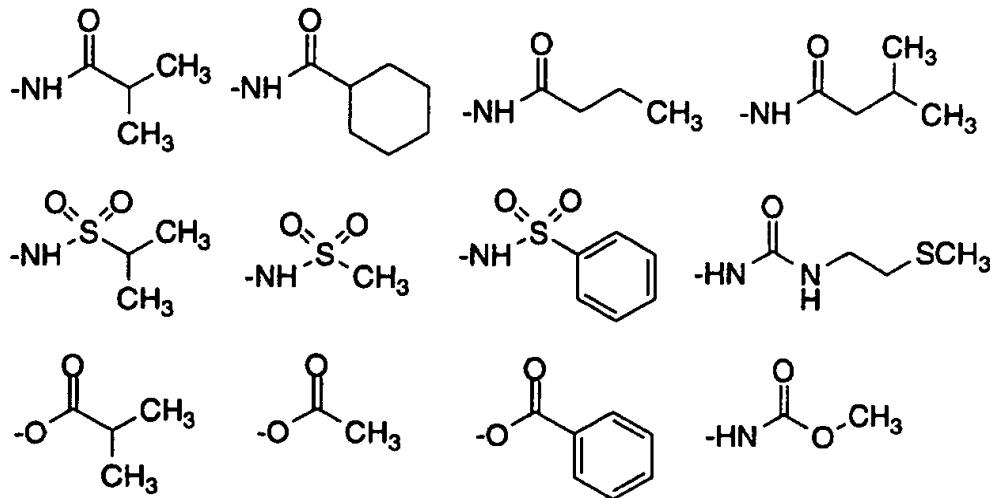
R^6 is selected from: hydrogen, C1-C6 alkyl, and $(CH_2)_v$ aryl, wherein the $(CH_2)_v$ and alkyl groups may be optionally substituted by $-O(R^2)$, $-S(O)_mR^2$, $-C(O)OR^2$, $-C(O)N(R^2)(R^2)$, $-SO_2N(R^2)(R^2)$, or

5 $-N(R^2)C(O)N(R^2)(R^2)$, wherein the aryl group is selected from: phenyl, pyridyl, 1H-tetrazolyl, triazolyl, oxadiazolyl, pyrazolyl, thiadiazoyl, and benzimidazol-2-yl, which is optionally substituted with C1-C6 alkyl, C3-C6 cycloalkyl, amino, or hydroxyl;

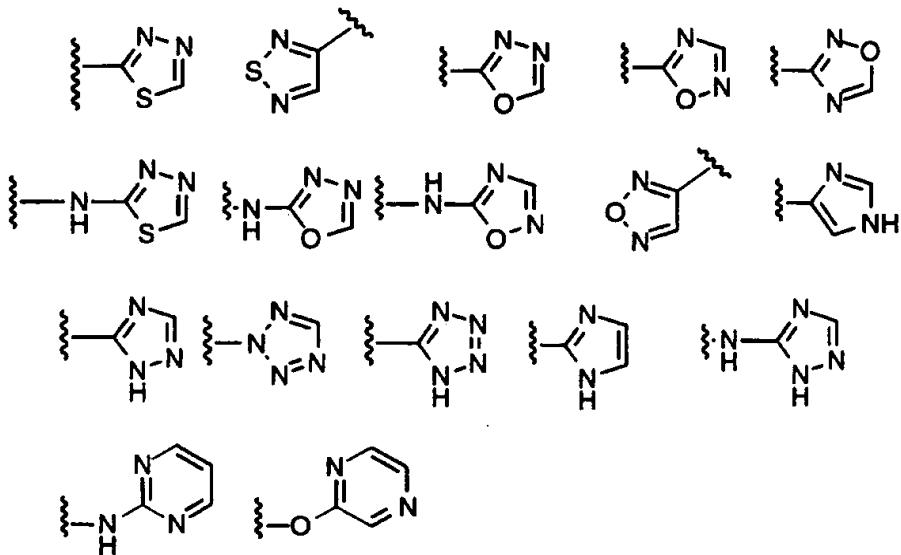
10 X is selected from the group consisting of: hydrogen,



- 18 -



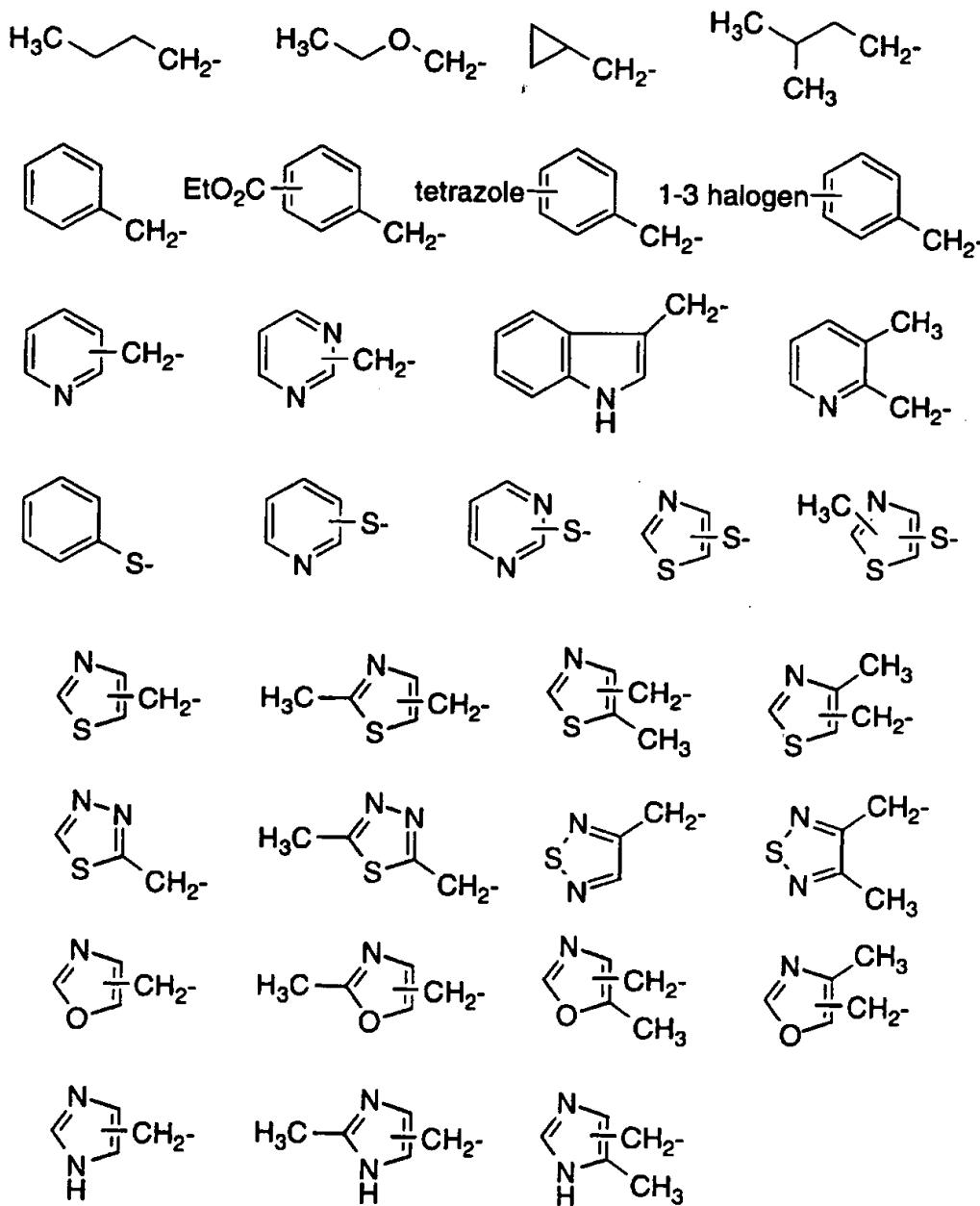
and further selected from the following group of heterocycles



5 wherein the heterocycle is optionally substituted with a substituent selected from: -N(R²)(R²), -O(R²), C₁-C₃ alkyl, halogen, and trifluoromethyl;

- 19 -

Y is selected from the group consisting of: hydrogen,



5 or their regioisomers whereof where not specified;

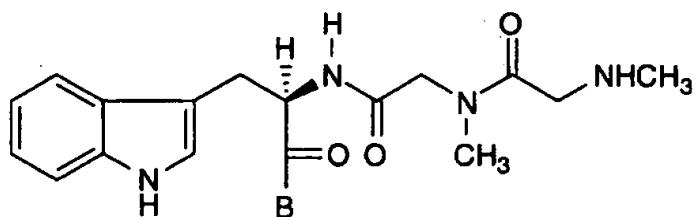
- 20 -

D is selected from: -N(R⁷)-, -S(O)_m-, -C(O)- and -C(H)(R⁷)-, wherein R⁷ is selected from: -R², -(CH₂)_qaryl, -C(O)R², -SO₂R², -C(O)N(R²)(R²), -C(O)OR², 1-H-tetrazol-5-yl, -SO₂N(R²)aryl, -SO₂N(R²)(R²) and the (CH₂)_q may be optionally substituted by C₁-C₄ alkyl, and the R² and aryl may be optionally further substituted with a substituent selected from: -OR^{2a}, -C(O)OR^{2a}, -C(O)N(R^{2a})(R^{2a}), halogen, -C₁-C₄ alkyl, and the aryl is selected from of triazolyl, oxadiazolyl, 1H-tetrazolyl, and thiadiazolyl;

5 10 1 is 0, 1 or 2;
m is 0, 1, or 2;
q is 0, 1, 2, 3, or 4;
r is 0, 1, 2, or 3;
t is 0, 1, 2, or 3;
15 v is 0, 1, or 2;

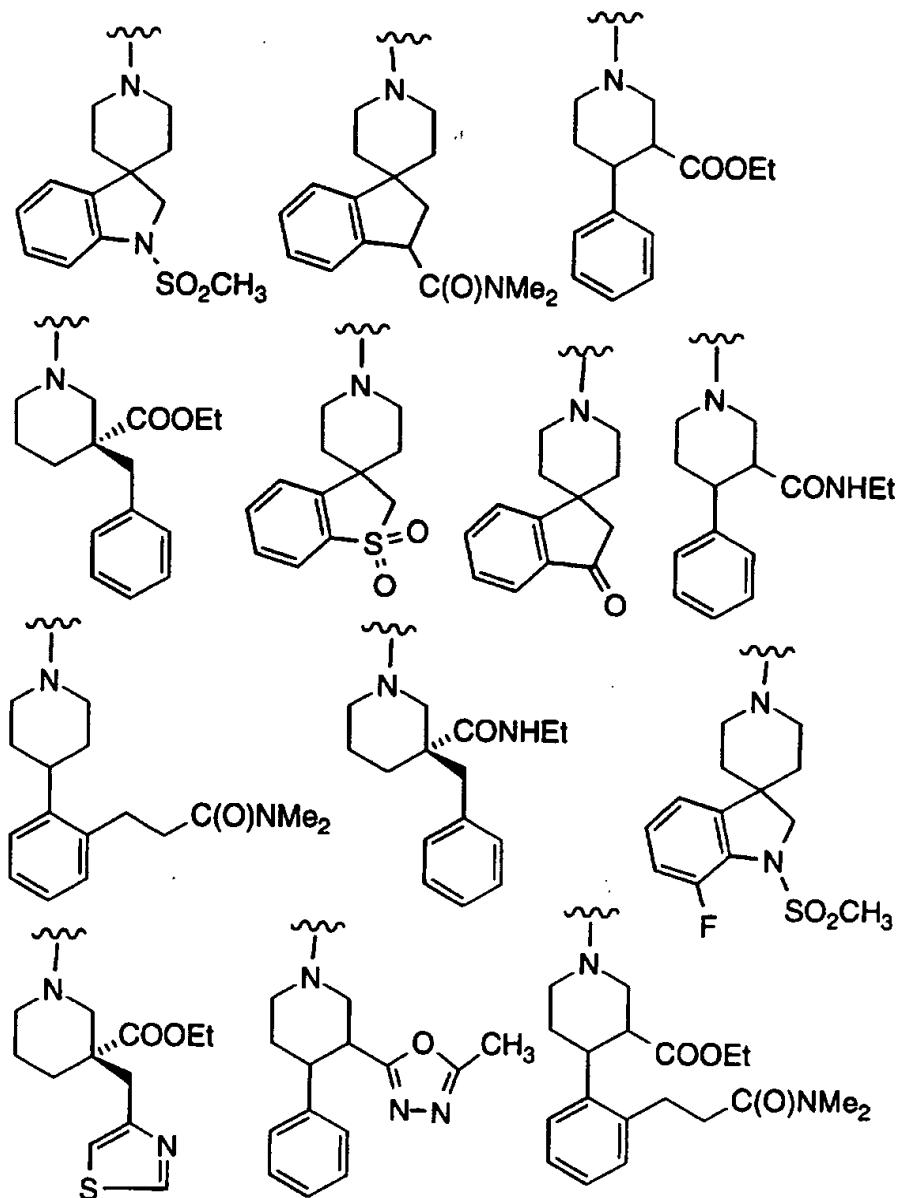
and pharmaceutically acceptable salts and individual diastereomers thereof.

20 One embodiment of the most preferred compounds of the instant invention includes compounds of the formula:



- 21 -

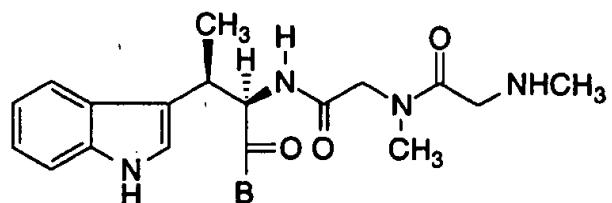
wherein B is selected from the group consisting of:



and pharmaceutically acceptable salts and individual diastereomers thereof.

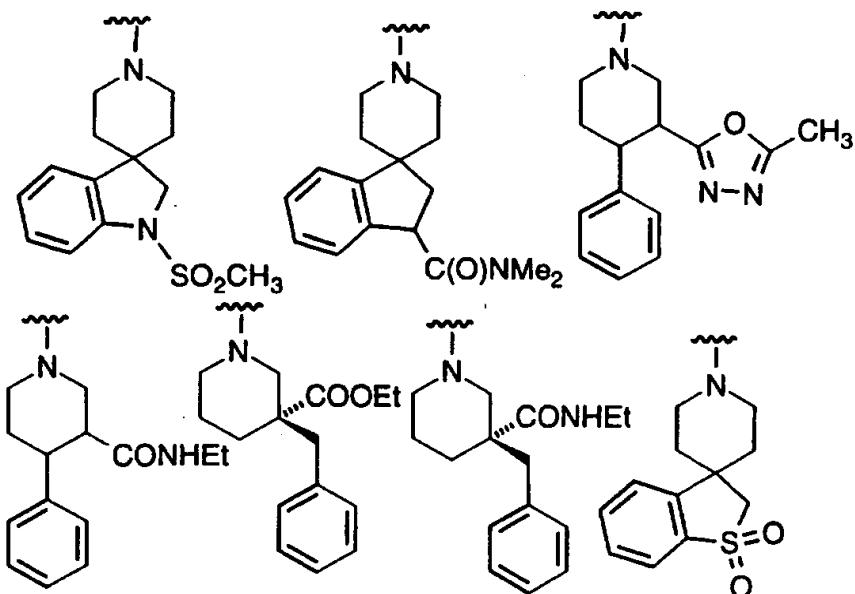
- 22 -

Another embodiment of the most preferred compounds of the instant invention includes compounds of the formula:



5

wherein B is selected from the group consisting of:

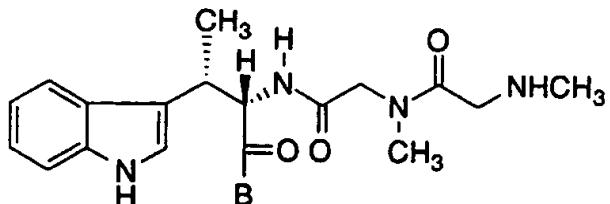


and pharmaceutically acceptable salts and individual diastereomers thereof.

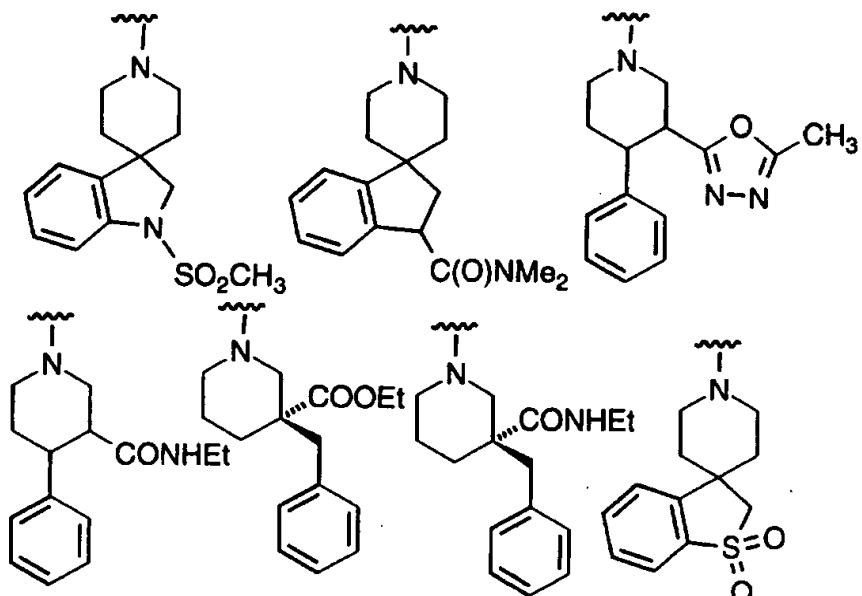
10

- 23 -

Another embodiment of the most preferred compounds of the instant invention includes compounds of the formula:



wherein B is selected from the group consisting of:

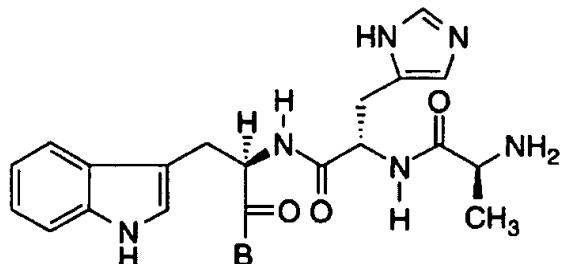


5

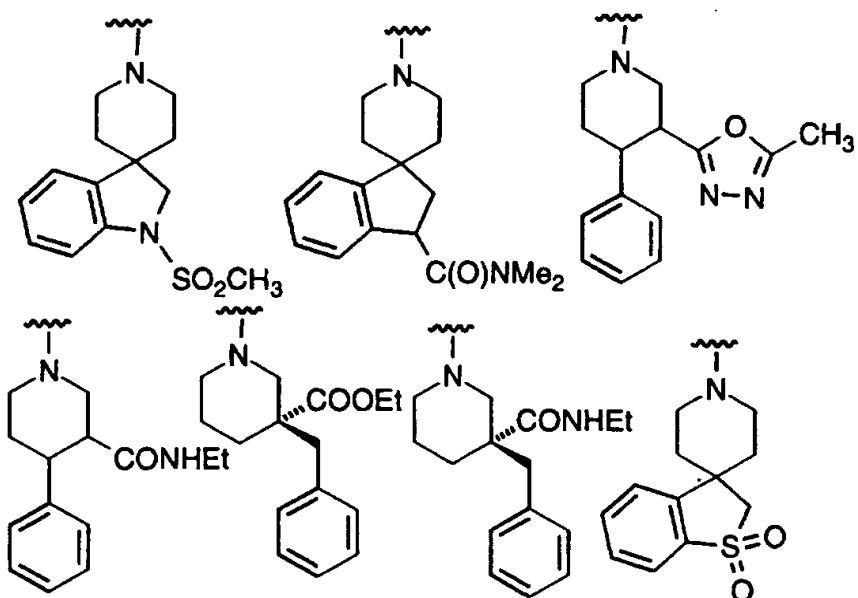
and pharmaceutically acceptable salts and individual diastereomers thereof.

- 24 -

Another embodiment of the most preferred compounds of the instant invention includes compounds of the formula:



wherein B is selected from the group consisting of:

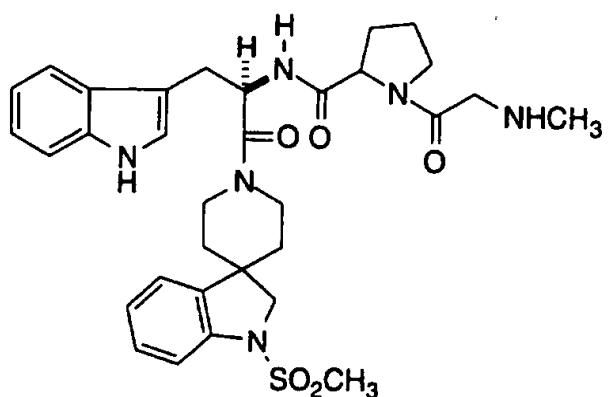
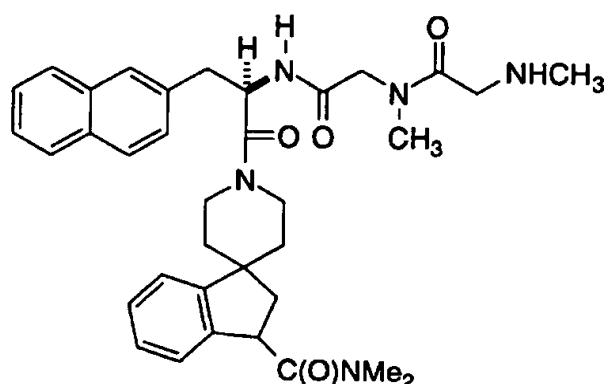
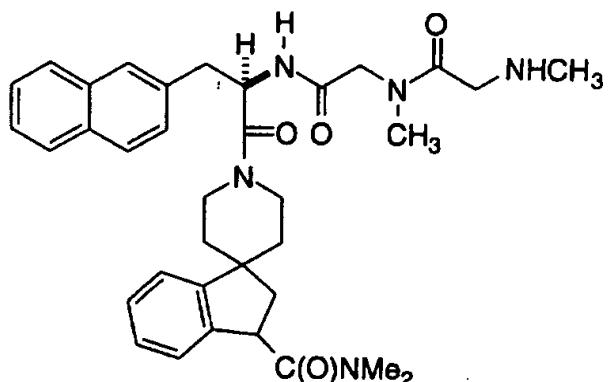


5

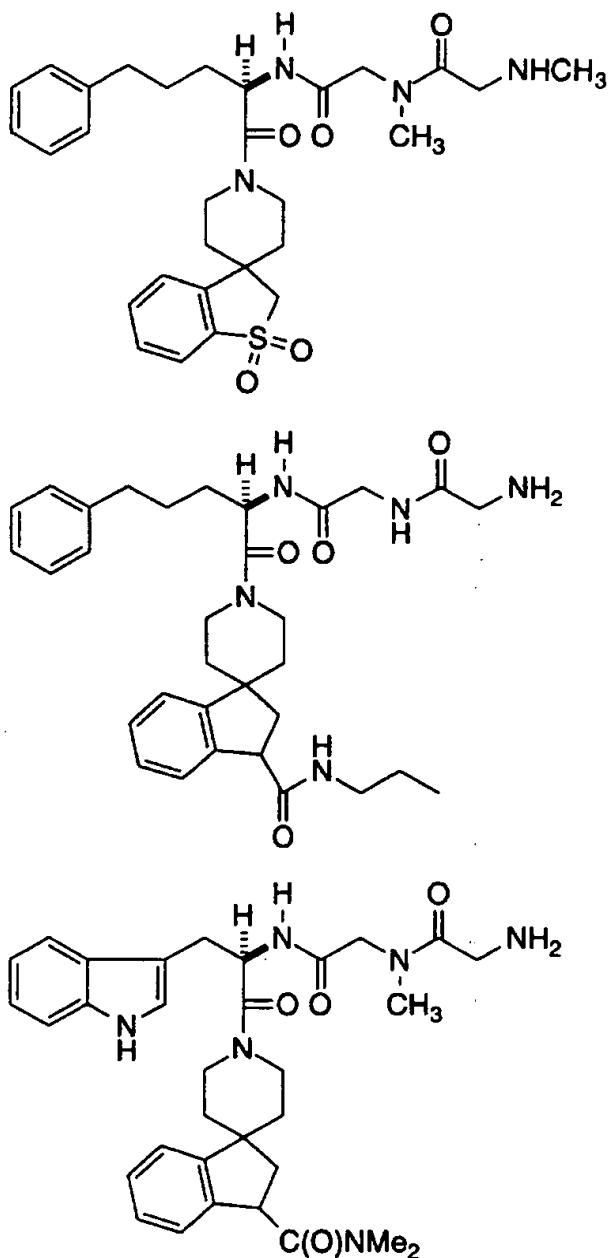
and pharmaceutically acceptable salts and individual diastereomers thereof.

- 25 -

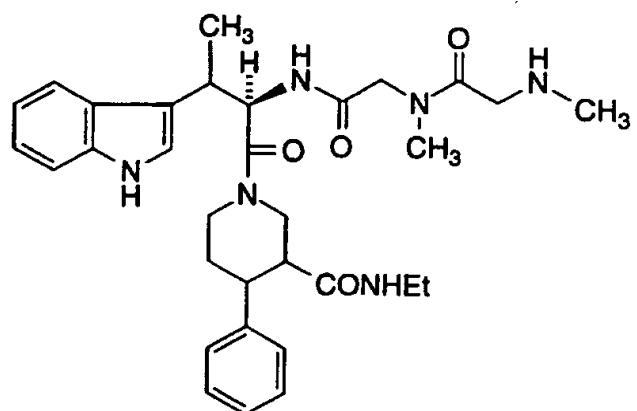
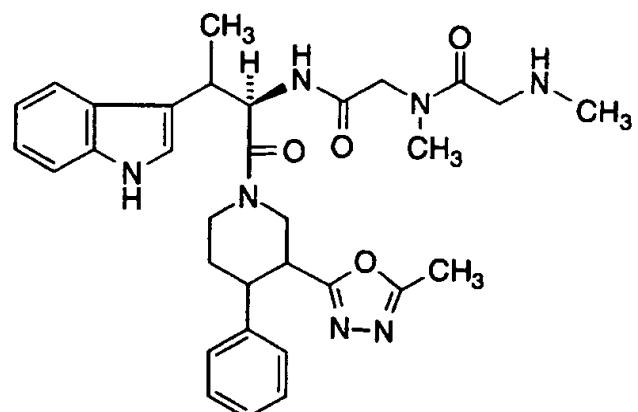
Specific compounds within the instant invention include the following:



- 26 -



- 27 -



and pharmaceutically acceptable salts and individual diastereomers
5 thereof where not otherwise specified.

- 28 -

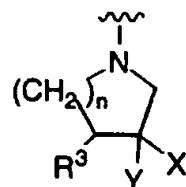
Throughout the instant application, the following abbreviations are used with the following meanings:

	Bu	butyl
	Bn	benzyl
5	BOC, Boc	t-butyloxycarbonyl
	BOP	Benzotriazol-1-yloxy tris(dimethylamino)- phosphonium hexafluorophosphate
	calc.	calculated
	CBZ, Cbz	Benzyloxycarbonyl
10	DCC	Dicyclohexylcarbodiimide
	DMF	N,N-dimethylformamide
	DMAP	4-Dimethylaminopyridine
	EDC	1-(3-dimethylaminopropyl)-3-ethylcarbodi-imide hydrochloride
15	EI-MS	Electron ion-mass spectroscopy
	Et	ethyl
	eq.	equivalent(s)
	FAB-MS	Fast atom bombardment-mass spectroscopy
	HOBT, HOBt	Hydroxybenztriazole
20	HPLC	High pressure liquid chromatography
	KHMDS	Potassium bis(trimethylsilyl)amide
	LAH	Lithium aluminum hydride
	LHMDS	Lithium bis(trimethylsilyl)amide
	Me	methyl
25	MF	Molecular formula
	MHz	Megahertz
	MPLC	Medium pressure liquid chromatography
	NMM	N-Methylmorpholine
	NMR	Nuclear Magnetic Resonance
30	Ph	phenyl
	Pr	propyl
	prep.	prepared
	TFA	Trifluoroacetic acid
	THF	Tetrahydrofuran

- 29 -

TLC Thin layer chromatography
 TMS Tetramethylsilane

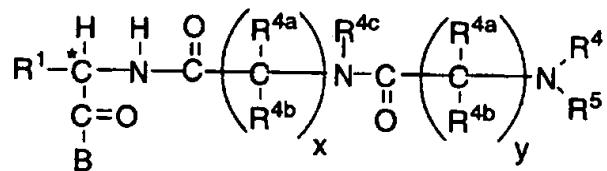
5 The compounds of the instant invention have at least
 two asymmetric centers when B is:



and both X and Y are groups other than hydrogen and are different from each other. Additional asymmetric centers may be present depending upon the nature of the various substituents on the molecule.

10 Each such asymmetric center will independently produce two optical isomers and it is intended that all of the possible optical isomers and diastereomers in mixture and as pure or partially purified compounds are included within the ambit of this invention. In the case of the asymmetric center which bears the X and Y groups, in most cases, both

15 R- and S- configurations are consistent with useful levels of growth hormone secretagogue activity. In addition configurations of many of the most preferred compounds of this invention are



20 Formula I

indicated. When the carbon atom in Formula I bearing an asterisk is of a defined two diastereomers result according to the absolute configuration at the carbon atom bearing the X and Y groups. These

25 diastereomers are arbitrarily referred to as diastereomer 1 (d1) and

- 30 -

diastereomer 2 (d2) in this invention and, if desired, their independent syntheses or chromatographic separations may be achieved as described herein. Their absolute stereochemistry may be determined by the x-ray crystallography of crystalline products or crystalline intermediates
5 which are derivatized, if necessary, with a reagent containing an asymmetric center of known absolute configuration.

The instant compounds are generally isolated in the form of their pharmaceutically acceptable acid addition salts, such as the salts derived from using inorganic and organic acids. Examples of such acids
10 are hydrochloric, nitric, sulfuric, phosphoric, formic, acetic, trifluoroacetic, propionic, maleic, succinic, malonic, methane sulfonic and the like. In addition, certain compounds containing an acidic function such as a carboxy can be isolated in the form of their inorganic salt in which the counterion can be selected from sodium, potassium,
15 lithium, calcium, magnesium and the like, as well as from organic bases.

The preparation of compounds of Formula I of the present invention may be carried out in sequential or convergent synthetic routes. Syntheses detailing the preparation of the compounds of Formula I in a sequential manner are presented in the following reaction
20 schemes.

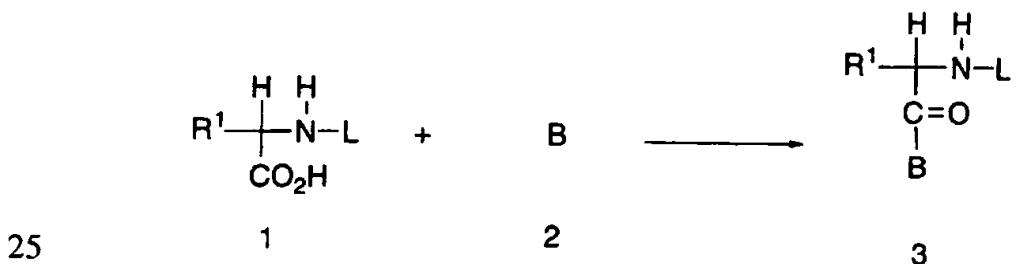
The phrase "standard peptide coupling reaction conditions" is used repeatedly here, and it means coupling a carboxylic acid with an amine using an acid activating agent such as EDC, DCC, and BOP in a inert solvent such as dichloromethane in the presence of a catalyst such
25 as HOBT. The uses of protective groups for amine and carboxylic acid to facilitate the desired reaction and minimize undesired reactions are well documented. Conditions required to remove protecting groups which may be present and can be found in Greene, T, and Wuts, P. G. M., *Protective Groups in Organic Synthesis*, John Wiley & Sons, Inc.,
30 New York, NY 1991. CBZ and BOC were used extensively in the synthesis, and their removal conditions are known to those skilled in the art. For example, removal of CBZ groups can be achieved by a number of methods known in the art; for example, catalytic hydrogenation with hydrogen in the presence of a nobel metal or its oxide such as palladium

- 31 -

on activated carbon in a protic solvent such as ethanol. In cases where catalytic hydrogenation is contraindicated by the presence of other potentially reactive functionality, removal of CBZ groups can also be achieved by treatment with a solution of hydrogen bromide in acetic 5 acid, or by treatment with a mixture of TFA and dimethylsulfide. Removal of BOC protecting groups is carried out in a solvent such as methylene chloride or methanol or ethyl acetate, with a strong acid, such as trifluoroacetic acid or hydrochloric acid or hydrogen chloride gas.

10 The protected amino acid derivatives 1 are, in many cases, commercially available, where the protecting group L is, for example, BOC or CBZ groups. Other protected amino acid derivatives 1 can be prepared by literature methods (Williams, R. M. *Synthesis of Optically Active α -Amino Acids*, Pergamon Press: Oxford, 1989). Many of the 15 piperidines, pyrrolidines, and hexahydro-1H-azepines of Formula 2 are either commercially available or known in the literature and others can be prepared following literature methods described for analogous compounds. Some of these methods are illustrated in the subsequent schemes. The skills required in carrying out the reaction and 20 purification of the resulting reaction products are known to those in the art. Purification procedures includes crystallization, normal phase or reverse phase chromatography.

SCHEME 1

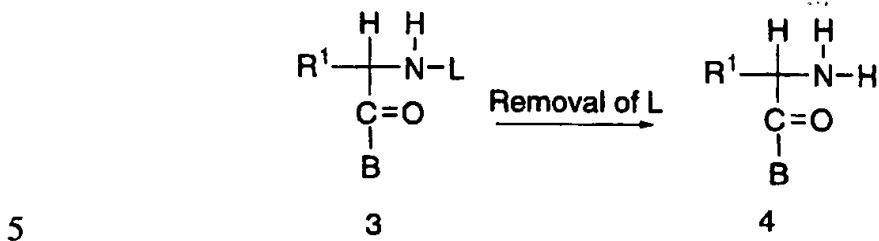


Intermediates of Formula 3 may be synthesized as described in Scheme 1. Coupling of amine of Formula 2, whose preparations are described later if they are not commercially available, to protected

- 32 -

amino acids of Formula 1, wherein L is a suitable protecting group, is conveniently carried out under standard peptide coupling conditions.

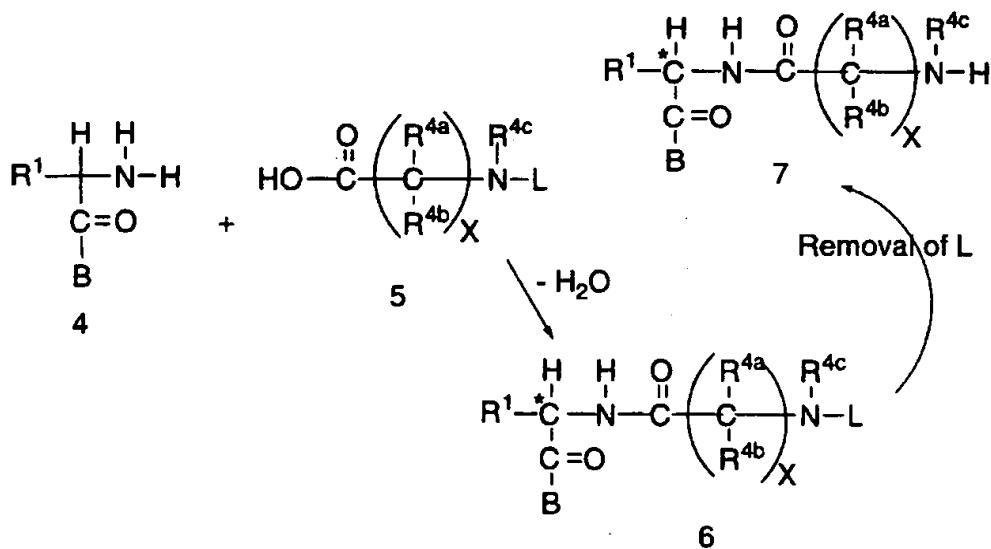
SCHEME 2



Conversion of 3 to intermediate 4 may be carried out as illustrated in Scheme 2 by removal of the protecting group L (CBZ, BOC, etc.)

10

SCHEME 3



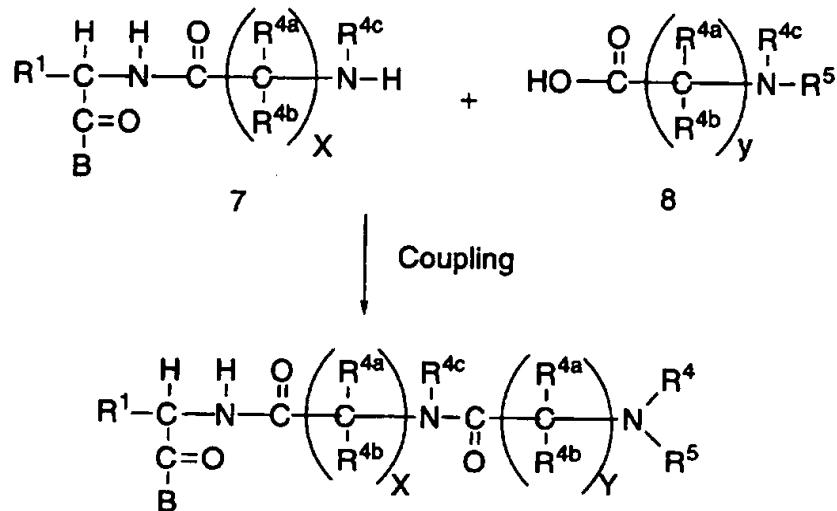
15 Intermediates of Formula 6, wherein L is a protecting group may be prepared as shown in Scheme 3 by coupling intermediates of Formula 4 to protected amino acids of Formula 5 under the standard peptide coupling reaction conditions. The amino acids 5, as amino acid 1, are either commercially available or can be synthesized by routine

- 33 -

methods. The removal of L in 6 to afford 7 can be carried out as noted above.

SCHEME 4

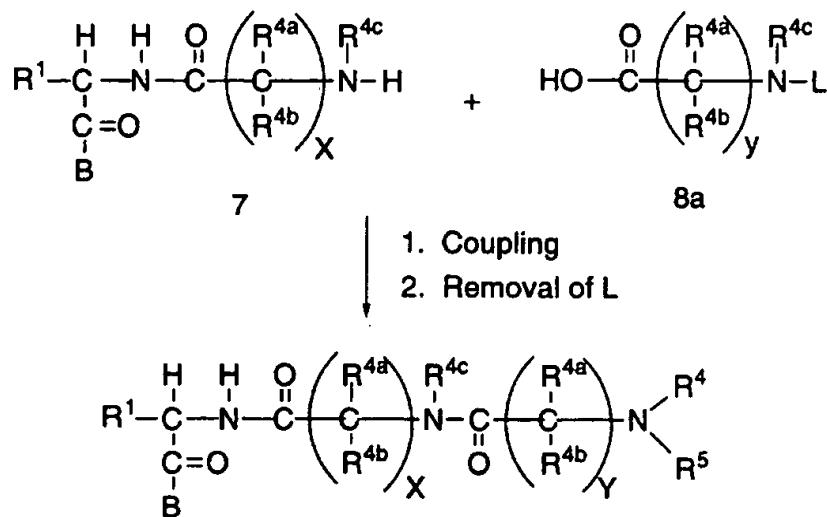
5



Elaboration of 7 to compounds of Formula 1 can be accomplished as shown in Scheme 4 by coupling intermediates of 10 Formula 7 to amino acids of Formula 8 under standard peptide coupling reaction conditions.

- 34 -

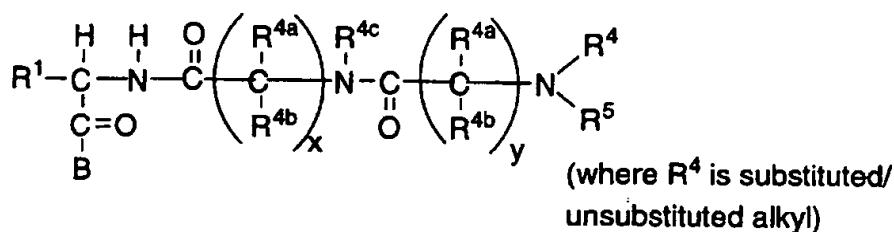
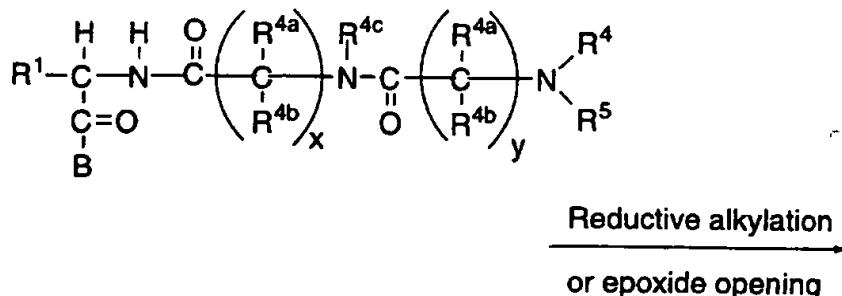
SCHEME 5



5 As shown in Scheme 5, if R^4 or R^5 is a hydrogen then the protected amino acids 8a are employed in the coupling reaction wherein L is a protecting group as defined above. The removal of L in 10 to afford I can be carried out as noted above.

- 35 -

SCHEME 6



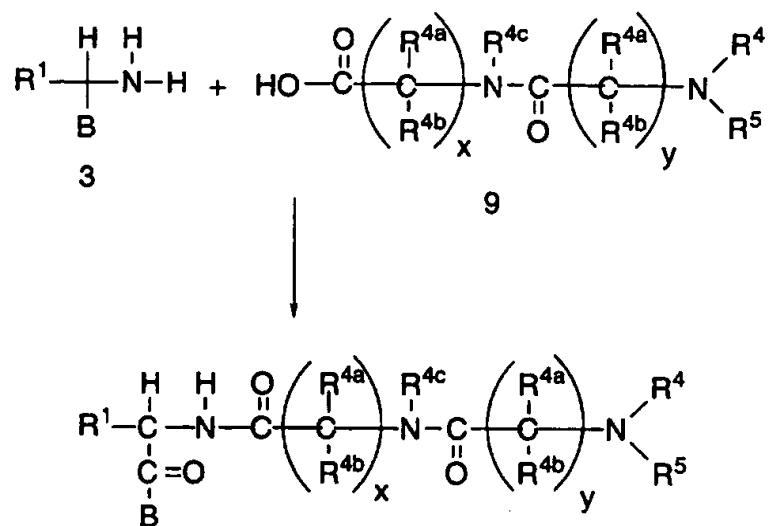
Compounds of Formula I wherein R⁴ and/or R⁵ is a

5 hydrogen may be further elaborated to new Compounds I which are substituted on the amino group as depicted in Scheme 6. Reductive alkylation of I with an aldehyde is carried out under conditions known in the art; for example, by catalytic hydrogenation with hydrogen in the presence of platinum, palladium, or nickel catalysts or with chemical 10 reducing agents such as sodium cyanoborohydride in a protic solvent such as methanol or ethanol in the present of catalytic amount of acid. Alternatively, a similar transformation can be accomplished via an epoxide opening reaction.

15 The compounds of general Formula I of the present invention may also be prepared in a convergent manner as described in Schemes 7, 8 and 9.

- 36 -

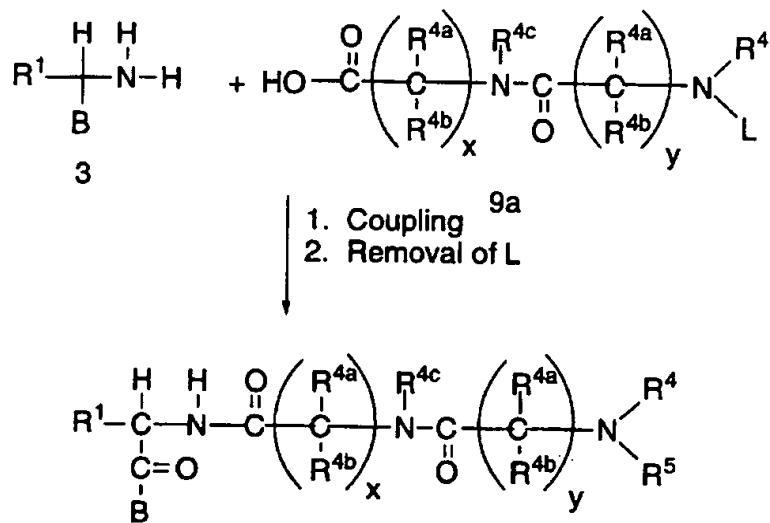
SCHEME 7



Intermediates of Formula 9 can be synthesized by well documented methods in the literature. Elaboration of 9 to compounds of Formula 1 can be accomplished as shown in Scheme 7 by coupling intermediates of Formula 3 to amino acids of Formula 9 under standard peptide coupling reaction conditions.

- 37 -

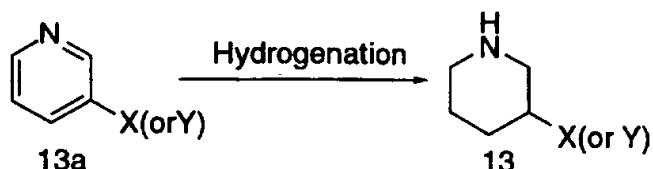
SCHEME 8



5 As shown in Scheme 8, if R⁴ or R⁵ is a hydrogen then the
 protected amino acids 9a are employed in the coupling reaction wherein
 L is a protecting group as defined above. The removal of L to afford I
 can be carried out as noted above. When R⁴ and/or R⁵ is H, substituted
 10 alkyl groups may be optionally added to the nitrogen atom as described
 in Scheme 4.

- 38 -

SCHEME 9

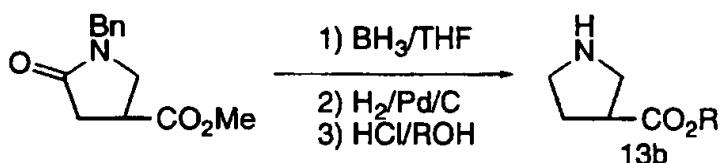


3-Monosubstituted piperidines of formula 13 can be

5 prepared by the reduction of pyridine derivatives or their salts by
hydrogenation in a suitable organic solvent such as water, acetic acid,
alcohol, e.g. ethanol, or their mixture, in the presence of a noble metal
catalyst such as platinum or an oxide thereof on a support such as
activated carbon, and conveniently at room temperature and
10 atmospheric pressure or under elevated temperature and pressure. 3-
Monosubstituted piperidines can also be prepared by modification of the
X or Y moiety of the existing 3-monosubstituted piperidines.

SCHEME 9A

15



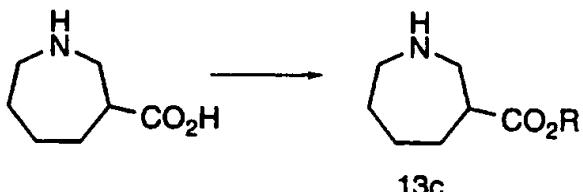
3-Monosubstituted pyrrolidines are commercially available
or can be conveniently prepared by literature procedures. Shown in
Scheme 9A is an example of the preparation of these compounds via
20 pyrrolidine-3-carboxylic acid ester. The commercially available
compound methyl 1-benzyl-4-oxo-3-pyrrolidinecarboxylate is reduced
by borane (*J. Chem. Soc.*, 24, 1618-1619). Removal of the benzyl
group by catalytic hydrogenolysis followed by ester exchange in an
appropriate alcohol medium such as ethyl alcohol in the presence of acid
25 gave the compound 13b. The ester functionality may be further
modified through conventional chemistry to other groups as defined by

- 39 -

X. 3-Monosubstituted pyrrolidines may also be prepared by catalytic hydrogenation of 3-substituted pyrroles.

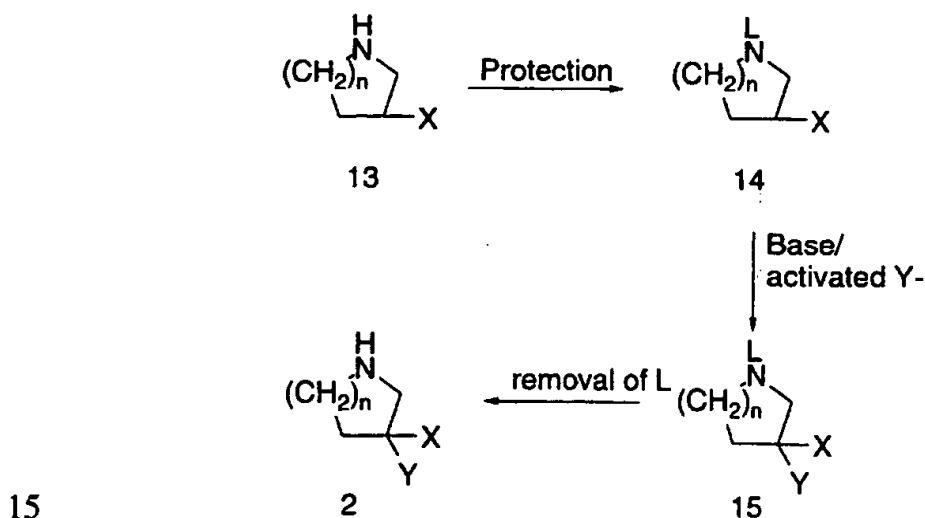
SCHEME 9B

5



Hexahydro-1H-azepines are commercially available or may be prepared by the literature procedure. Hexahydro-1H-azepine-3-carboxylic acid (Krogsgaard-Larsen, P. et al., Acta. Chem. Scand., **B32**, 327, (1978)) is esterified in an alcohol solvent in the presence of acid. The ester functionality may be further modified through conventional chemistry to other groups within the definition of X.

SCHEME 10



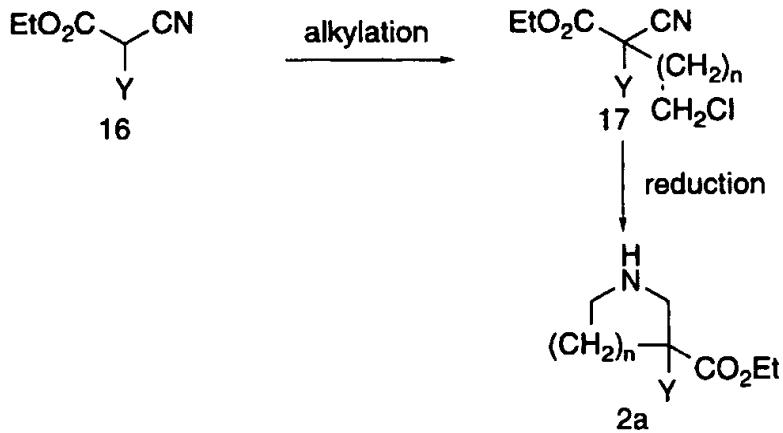
Illustrated in Scheme 10 is a general way to prepare di-substituted piperidines, pyrrolidines, and hexahydro-1H-azepines. Compounds of Formula 13 wherein X is an electron withdrawing group such as -CN, -CO2Rg, where Rg is alkyl, aryl, and (C1-C4alkyl)aryl are

- 40 -

known compounds or may be prepared by methods analogous to those used for the preparation of such known compounds. The secondary amine of compounds of Formula 13 may be first protected by a protecting group L such as BOC and CBZ using the conventional 5 techniques. Introduction of the Y substitution can be achieved by first reacting compounds of Formula 14 with a strong base such as lithium bis(trimethylsilyl)amide, lithium diisopropylamide following by addition of alkylating or acylating reagents such as alkyl halides, aryl alkyl halides, acyl halides, and haloformates in a inert solvent such as 10 THF at temperatures from -100° to room temperature. Thio derivatives where the sulfur is attached directly to an alkyl or an aryl group can be prepared similarly by reacting with a disulfide. The halides used in these reactions are either commercially available or known compounds in the literature or may be prepared by methods analogous to those used 15 for the preparation of known compounds. The protecting group L in compounds of formula 15 may be removed with conventional chemistry to give compounds of Formula 2.

SCHEME 11

20

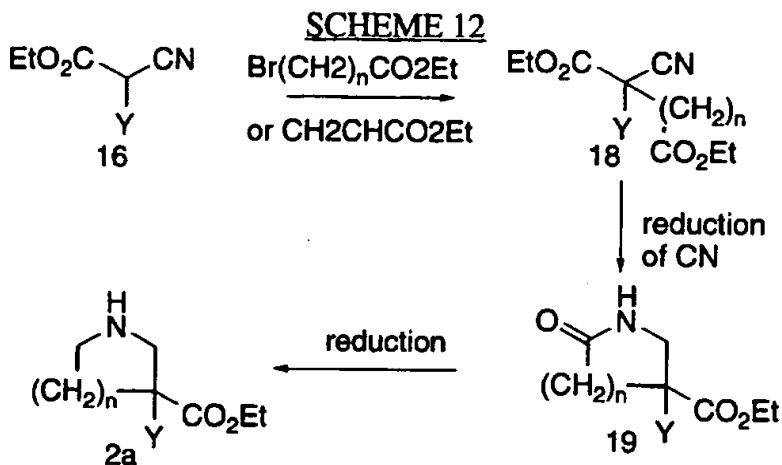


Alternative ways of preparing compounds of Formula 2 include construction of the ring itself (Jacoby, R. L. et al, J. Med. Chem., 17, 453-455, (1974)). Alkylation of the cyanoacetates of 25 general formula 16, which are commercially available or may be prepared from literature procedures, with alkyl dihalides such as 1-

- 41 -

bromo-2-chloroethane or 1-bromo-3-chloropropane yields the chloride 17. Reduction of the nitriles 17 by borane or by hydrogenation using Raney Ni as a catalyst gives the corresponding primary amines, which upon refluxing in ethanol to give compounds of Formula 2a.

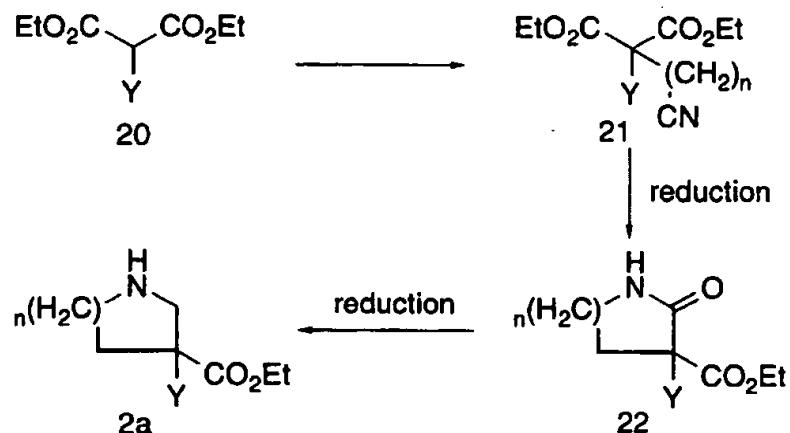
5



Alternatively, the cyanoacetates of general formula 16 may be alkylated with an ethoxycarbonylalkyl bromide or reacted with ethyl acrylate to give compounds of Formula 18. Reduction of the nitriles 18 by borane or by hydrogenation using Raney Ni as a catalyst gives the corresponding primary amines, which upon refluxing in ethanol gives lactam 19. Reduction of the lactam 19 by borane gives compounds of Formula 2a.

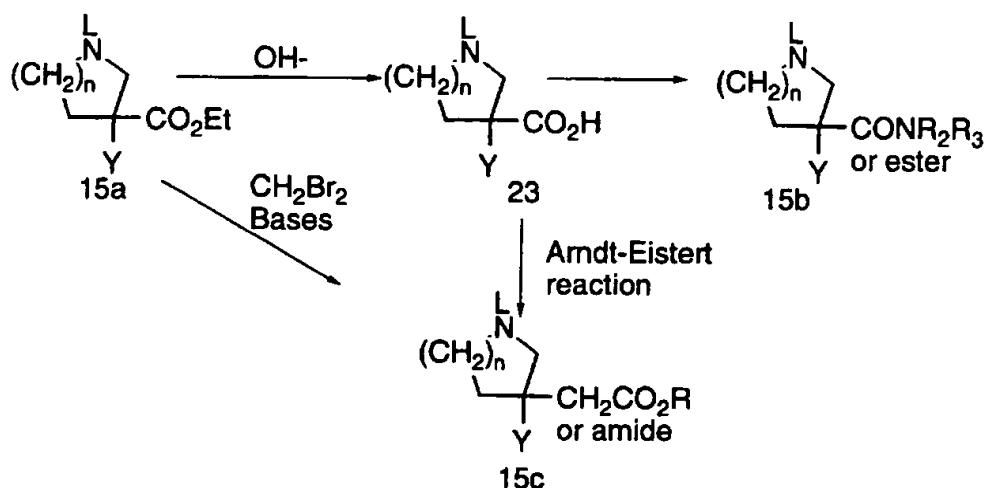
15

- 42 -

SCHEME 13

5 Alternatively, a malonate of general formula 20 may be
 alkylated with cyanoalkyl bromide or can be reacted with acrylonitrile
 to form compounds of formula 21. Reduction of the nitriles 21 by
 borane or by hydrogenation using Raney Ni as a catalyst gives the
 corresponding primary amines, which upon refluxing in ethanol gives
 10 lactam 22. Reduction of the lactam 22 by borane gives compounds of
 formula 2a.

SCHEME 14

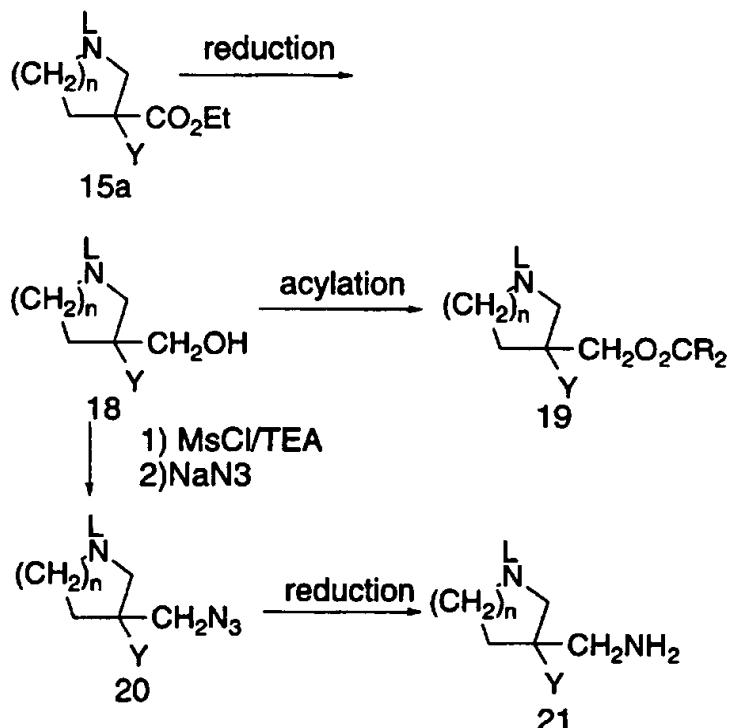


The X, Y functionalities in compounds of general structure

5 15 may be further elaborated to groups not accessible by direct
alkylation. For example in Compound 15 when $X = \text{CO}_2\text{Et}$ the ester
(provided that this is the only ester group in the molecule) can be
saponified to the carboxylic acid, which can be further derivatized to
amides or other esters. The carboxylic acid can be converted into its
10 next higher homologue, or to a derivative of the homologous acid, such
as amide or ester by an Arndt-Eistert reaction. Alternatively, the ester
can be directly homologated by the protocol using ynone anions
described by C. J. Kowalski and R. E. Reddy in J. Org. Chem., 57,
15 7194-7208 (1992). The resulting acid and/or ester may be converted to
the next higher homologue, and so on and so forth. The protecting
group L may be removed through conventional chemistry.

- 44 -

SCHEME 15

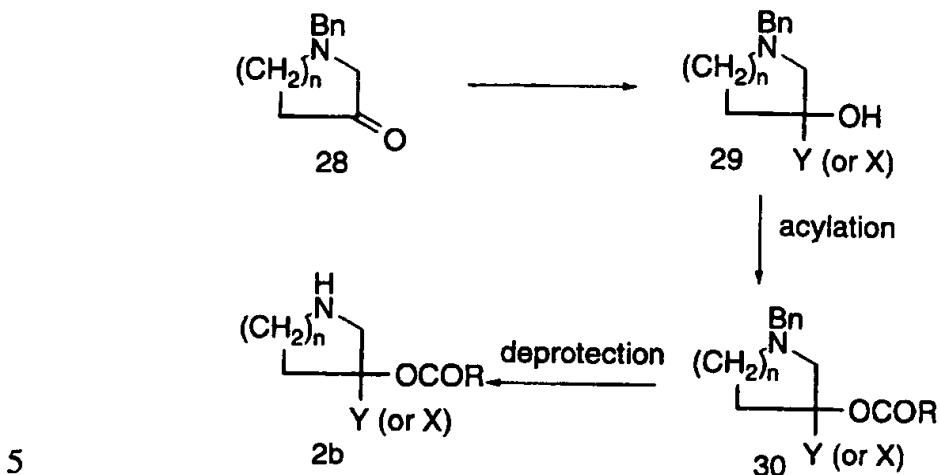


The ester in 15a may be reduced to an alcohol 18 in a suitable solvent such as THF or ether with a reducing agent such as DIBAL-H and conveniently carried out at temperatures from -100°C to 0°C. The alcohol may be acylated to Compound 19 in a suitable solvent such as dichloromethane using an acyl halide or acid anhydride in the presence of a base such as triethyl amine (TEA). The hydroxy group in 18 may also be converted to a good leaving group such as mesylate and displaced by a nucleophile such as cyanide, a thiol or an azide.

Reduction of the azide in compounds of Formula 20 to an amine 21 can be achieved by hydrogenation in the presence of a noble metal such as palladium or its oxide or Raney nickel in a protic solvent such as ethanol. The nitrile can be reduced to afford the homologous amine. The amine of Formula 21 may be further elaborated to amides, ureas sulfonamides as defined by X through conventional chemistry. The protecting group L may be removed through conventional chemistry.

- 45 -

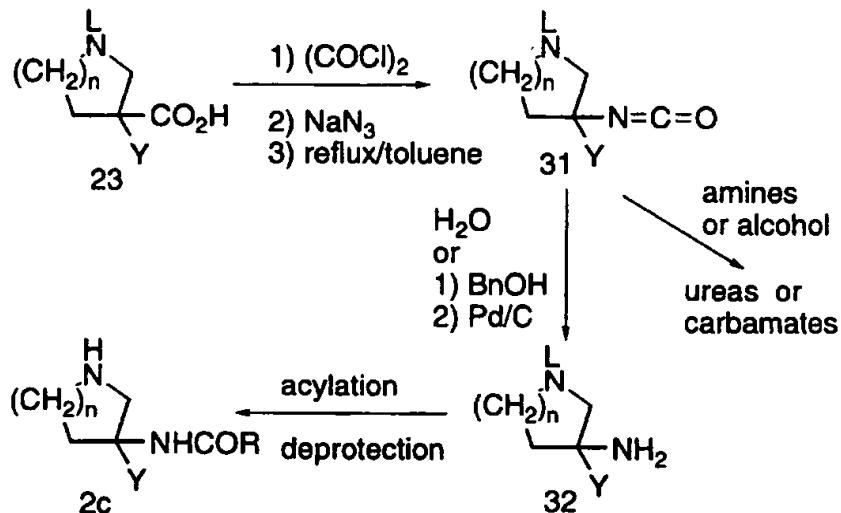
SCHEME 16



5 In cases where oxygen is directly attached to the ring, a convenient method involves the addition reaction by an activated form of an alkyl, aryl, alkylaryl group, such as lithium reagent, Grignard reagents, and the like with a ketone of general formula 28, which is
10 commercially available. Further derivatization of the resulting hydroxy group by acylation, sulfonylation, alkylation, and the like gives compounds as defined by Y or X through conventional chemistry.
15 Removal of the benzyl protective group may be carried out under the usual conditions to give compounds of general formula 2b. Shown in Scheme 16 is a general example of acylations.

- 46 -

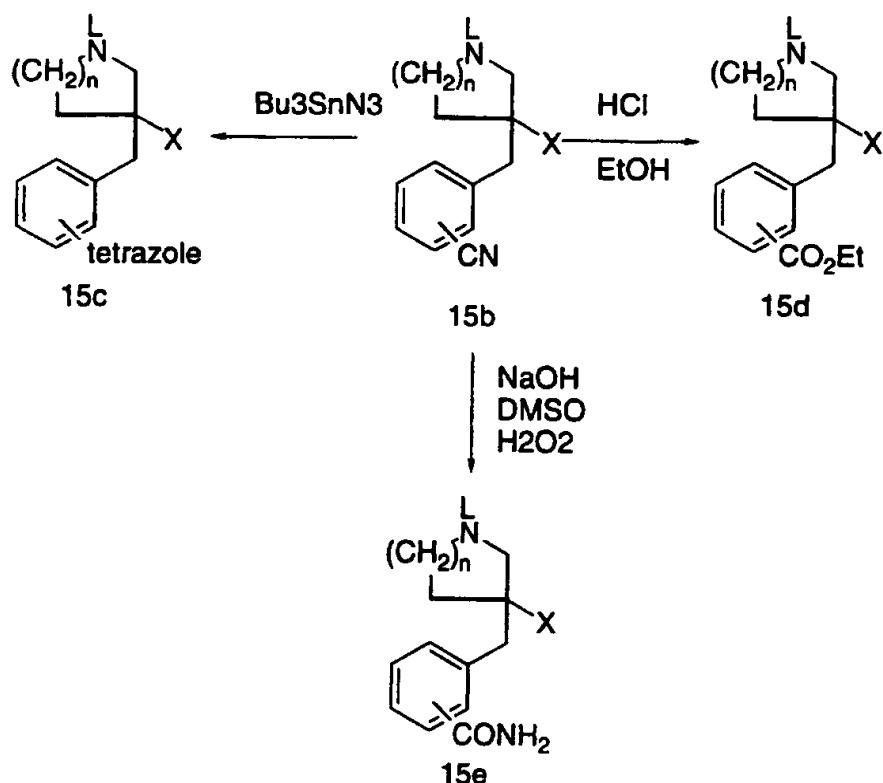
SCHEME 17



In cases where a nitrogen-substituted group is directly attached to the ring, a convenient method is to use the Curtius rearrangement on the acid 23 to afford the isocyanate 31. Addition of amines or alcohols give ureas or carbamates respectively which can be deprotected to remove L to give special cases of compounds of formula 2. Conversion of the isocyanate to amine by hydrolysis gives compound 32. Further derivatization of the resulting amine group by acylation, sulfonylation, alkylation, and the like to give compounds as defined by Y or X can be done through conventional chemistry. Removal of the protective group L may be carried out under the usual conditions to give compounds of general formula 2c. Shown in Scheme 17 is a general example of acylations.

- 47 -

SCHEME 18



For compounds that are not readily obtainable by direct

5 alkylation as shown in Scheme 10, modifications of easily obtainable
 compounds of general formula 15 may be conducted to achieve the
 desired substitution through conventional chemistry. For example,
 compounds with Y being hydroxybenzyl may be prepared by
 demethylation of the corresponding compound wherein Y is

10 methoxybenzyl. Similarly, compounds with Y being aminobenzyl may
 be prepared by reduction of the corresponding compound wherein Y is
 nitrobenzyl. Shown in Scheme 18 is an example of a procedure that
 uses nitrile as a starting point for the preparation of compounds with
 different substitutions. Removal of the protective group L gives
 15 compounds of general formula 2 as described in Scheme 10.

Compounds of the general formula 2 prepared in this way
 are racemic when X and Y are not identical. Resolution of the two
 enantiomers can be conveniently achieved by classical crystallization

- 48 -

methods by using a chiral acid such as L- or D-tartaric acid, (+) or (-) 10-camphorsulfonic acid in a suitable solvent such as acetone, water, alcohol, ether, acetate or their mixture. Alternatively, the racemic amine 2 can be reacted with a chiral auxiliary such as (R) or (S)-O-

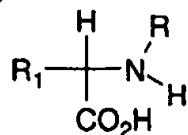
5 acetylmandelic acid followed by chromatographic separation of the two diastereomers, and removal of the chiral auxiliary by hydrolysis. Alternatively asymmetric alkylation can also be utilized for the synthesis of optically active intermediate by introducing a removable chiral auxiliary in X or in place of L with subsequent chromatographic

10 separation of diastereomers.

In cases where a sulfide is present in the molecule, it may be oxidized to a sulfoxide or to a sulfone with oxidizing agents such as sodium periodate, m-chloroperbenzoic acid or Oxone® in an solvent such as dichloromethane, alcohol or water or their mixtures.

15 The compounds of the present invention may also be prepared from a variety of substituted natural and unnatural amino acids of formulas 46. The preparation of many of these acids is described in US Patent No. 5,206,237. The preparation of these intermediates in racemic form is accomplished by classical methods familiar to those skilled in the art (Williams, R. M. "Synthesis of Optically Active α -Amino Acids" Pergamon Press: Oxford, 1989; Vol. 7). Several methods exist to resolve (DL)-

20



46

amino acids. One of the common methods is to resolve amino or carboxyl protected intermediates by crystallization of salts derived from optically active acids or amines. Alternatively, the amino group of carboxyl protected intermediates may be coupled to optically active acids by using chemistry described earlier. Separation of the individual diastereomers either by chromatographic techniques or by

25 crystallization followed by hydrolysis of the chiral amide furnishes resolved amino acids. Similarly, amino protected intermediates may be

30

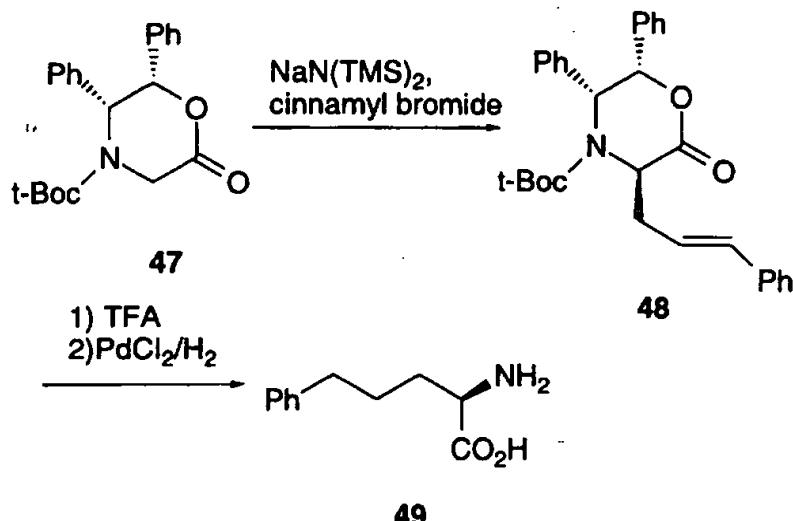
- 49 -

converted to a mixture of chiral diastereomeric esters and amides. Separation of the mixture using methods described above and hydrolysis of the individual diastereomers provides (D) and (L) amino acids. Finally, an enzymatic method to resolve N-acetyl derivatives of (DL)-
5 amino acids has been reported by Whitesides and coworkers in *J. Am. Chem. Soc.* 1989, 111, 6354-6364.

When it is desirable to synthesize these intermediates in optically pure form, established methods include: (1) asymmetric electrophilic amination of chiral enolates (*J. Am. Chem. Soc.* 1986, 108,
10 6394-6395, 6395-6397, and 6397-6399), (2) asymmetric nucleophilic amination of optically active carbonyl derivatives, (*J. Am. Chem. Soc.* 1992, 114, 1906; *Tetrahedron Lett.* 1987, 28, 32), (3) diastereoselective alkylation of chiral glycine enolate synthons (*J. Am. Chem. Soc.* 1991, 113, 9276; *J. Org. Chem.* 1989, 54, 3916), (4) diastereoselective
15 nucleophilic addition to a chiral electrophilic glycinate synthon (*J. Am. Chem. Soc.* 1986, 108, 1103), (5) asymmetric hydrogenation of prochiral dehydroamino acid derivatives ("Asymmetric Synthesis, Chiral Catalysis; Morrison, J. D., Ed; Academic Press: Orlando, FL, 1985; Vol 5), and (6) enzymatic syntheses (*Angew. Chem. Int. Ed. Engl.* 1978, 17, 176).

- 50 -

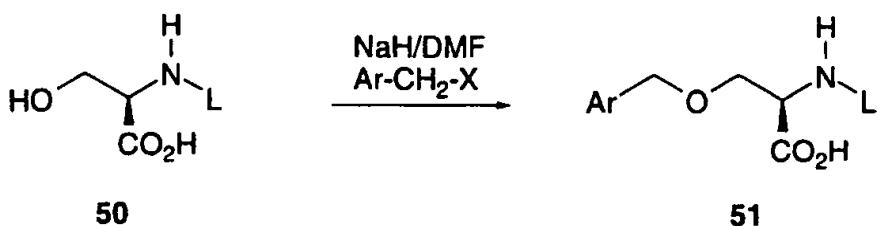
SCHEME 19



For example, alkylation of the enolate of diphenyloxazinone 47 (*J. Am. Chem. Soc.* 1991, **113**, 9276) with cinnamyl bromide in the presence of sodium bis(trimethylsilyl)amide proceeds smoothly to afford 48 which is converted into the desired (D)-2-amino-5-phenylpentanoic acid 49 by removing the N-t-butyloxycarbonyl group with trifluoroacetic acid and hydrogenation over a PdCl_2 catalyst (Scheme 19).

10

SCHEME 20



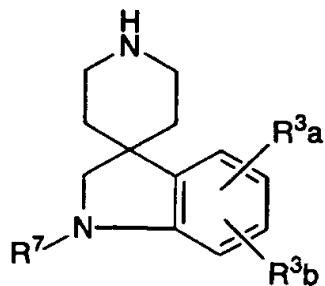
15 Intermediates of formula 46 which are O-benzyl-(D)-serine derivatives 51 are conveniently prepared from suitably substituted benzyl halides and N-protected-(D)-serine 50. The protecting group L is conveniently a BOC or a CBZ group. Benzylation of 64 can be achieved by a number of methods well known in the literature including

- 51 -

deprotonation with two equivalents of sodium hydride in an inert solvent such as DMF followed by treatment with one equivalent of a variety of benzyl halides (*Synthesis* 1989, 36) as shown in Scheme 20.

The O-alkyl-(D)-serine derivatives may also be prepared 5 using an alkylation protocol. Other methods that could be utilized to prepare (D)-serine derivatives of formula 51 include the acid catalyzed benzylation of carboxyl protected intermediates derived from 50 with reagents of formula $\text{ArCH}_2\text{OC}(=\text{NH})\text{CCl}_3$ (O. Yonemitsu *et al.*, *Chem. Pharm. Bull.* 1988, 36, 4244). Alternatively, alkylation of the chiral 10 glycine enolates (*J. Am. Chem. Soc.* 1991, 113, 9276; *J. Org. Chem.* 1989, 54, 3916) with $\text{ArCH}_2\text{OCH}_2\text{X}$ where X is a leaving group affords 51. In addition D,L-O-aryl(alkyl)serines may be prepared and resolved 15 by methods described above.

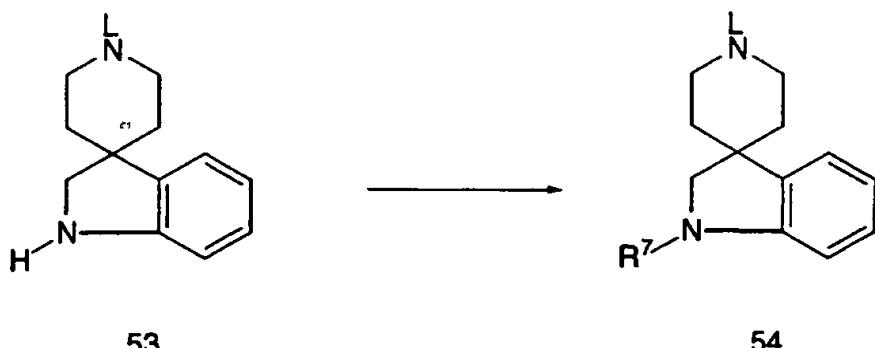
The spiro piperidines of formula 52 may be prepared by a 15 number of methods, including the syntheses described below.



52

- 52 -

SCHEME 21

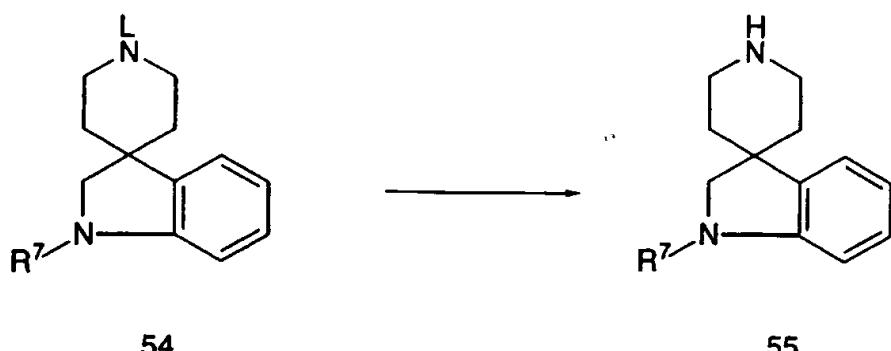


As shown in Scheme 21, the spiropiperidine of formula 43, wherein L is a defined protecting group, is synthesized by methods that

5 are known in the literature (for example H. Ong et al *J. Med. Chem.* 1983, 23, 981-986). The indoline nitrogen of 54, wherein L is a protecting group such as methyl or benzyl, can be reacted by with a variety of electrophiles to yield spiro piperidines of formula 54, wherein R⁹ can be a variety of functionalities. Compound 54 can be reacted with, for example, isocyanates in an inert solvent like dichloromethane to yield urea derivatives, chloroformates in an inert solvent such as dichloromethane to yield carbamates, acid chlorides, anhydrides, or acyl imidazoles to generate amides, sulfonyl chlorides to generate sulfonamides, sulfamyl chlorides to yield sulfamides. Also, the 10 indoline nitrogen of 53 can be reductively alkylated with aldehydes with conditions known in the art. When the aldehyde used in the reductive amination reaction is a protected glyoxylic acid of structure 15 HCOCOOM, wherein M is a defined protecting group, M can be removed from the product and further derivatized. Alternatively, 53 can be reacted with epoxides to produce 53, wherein R⁹ is β -hydroxy- 20 substituted alkyl or arylalkyl groups. The indoline 53 can also be transformed to compounds of formula 54, wherein R⁹ = phenyl or substituted phenyl, heteroaryl or substituted heteroaryl, by carrying out the reacting 53 with a fluoro phenyl or fluoro heteroaryl reagent. This 25 chemistry is detailed by H. Ong et al *J. Med. Chem.* 1983, 23, 981-986.

- 53 -

SCHEME 22

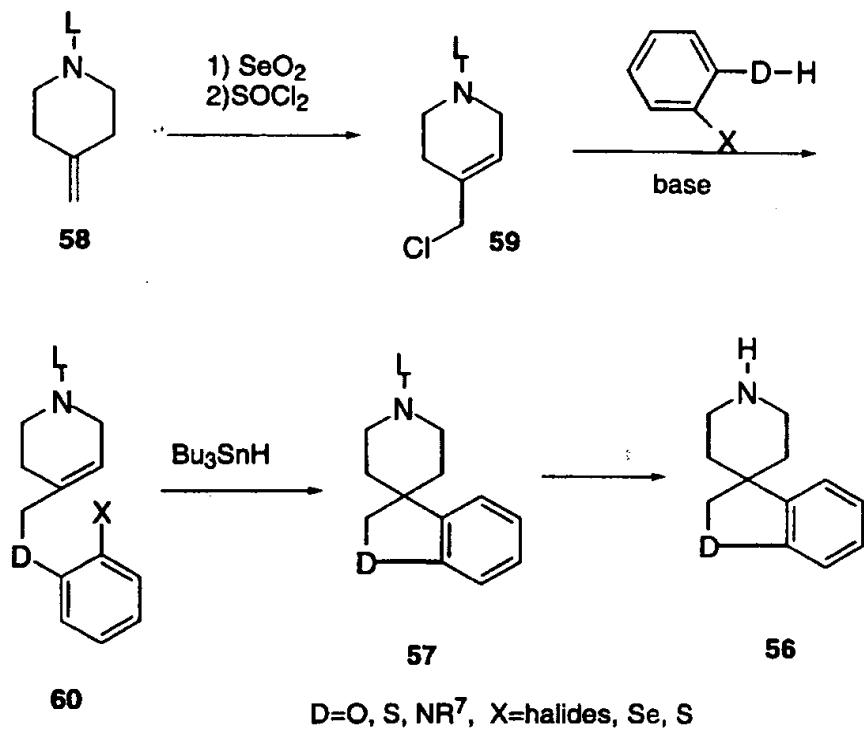


The spiro piperidine intermediate 54 (L = Me or Bn), wherein R⁷ is hydrogen or most of the derivatives described above, can be demethylated or debenzylated to produce 55, wherein R⁹ is hydrogen or most of the derivatives described above, as shown in Scheme 22. For compounds of formula 54, wherein L = Me, demethylation can be carried out by a number methods familiar to those skilled in the art. For example, demethylation of 54 be accomplished by reacting it with cyanogen bromide and potassium carbonate in an inert solvent solvent such as dichloromethane to yield a cyanamide which can reduced to give 55 by treatment with lithium aluminum hydride in refluxing tetrahydrofuran, refluxing strong acid like aqueous hydrochloric acid, or with Grignard reagents like methyl magnesium bromide.

Alternatively, demethylation of 54 can be effected with the ACE-Cl method as described in R. Olofson et al. *J. Org. Chem.* 1984, 49, 2795 and references therein. For intermediates of formula 54, wherein L = Bn, removal of benzyl group can be accomplished by reductive methods including hydrogenation in the presence of platinum or palladium catalyst in a protic solvent like methanol. Alternatively, debenzylation of 54, L = Bn, can be effected with the ACE-Cl method as described in R. Olofson et al. *J. Org. Chem.* 1984, 49, 2795 and references therein.

- 54 -

SCHEME 23

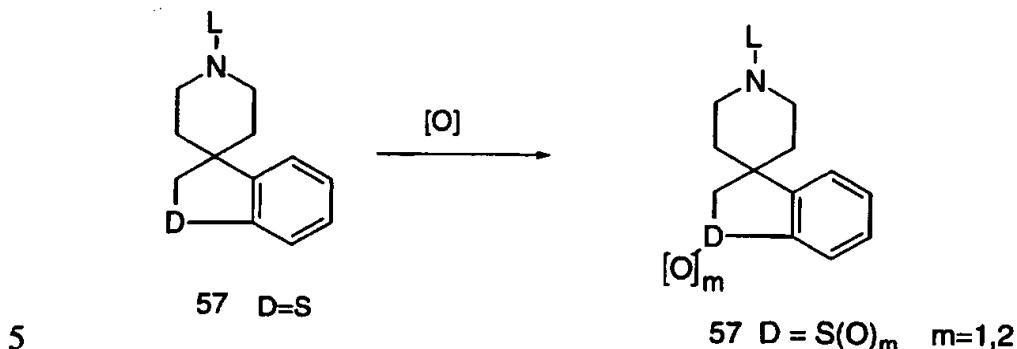


The spiro heterocyclic compounds of formula 56 can be prepared by a number of methods, including the syntheses as described in Scheme 23. Allylic oxidation of the protected piperidine 58 is accomplished by classical methods familiar to those skilled in the art (Rabjohn, N. *Org. React.* 1976, 24, 261). The resulting allylic alcohol is treated with thionyl chloride in an inert solvent such as benzene to provide the corresponding chloride 59. When $\text{D}=\text{O}$ or S , the alkylation is carried out in DMF or acetone as solvent with potassium carbonate as a base, and when $\text{D}=\text{NR}^7$ ($\text{R}^7=\text{H, alkyl, aryl, acyl, sulfonyl, carbamate}$) the reaction is carried out with sodium hydride as a base in an inert solvent such as THF to afford the cyclization precursor 60. When L is a defined protecting group, compound 60 can be cyclized by a number of methods familiar to those skilled in the art. For example, cyclization of 60 can be accomplished by reaction with tributyltin hydride (Curran, D. P. *Synthesis* 1988, 417 and 489) in an inert solvent such as benzene.

- 55 -

to yield 57. Alternatively, compound 57 (D=NR₉) can be prepared by the method shown in Schemes 24 and 25.

SCHEME 24



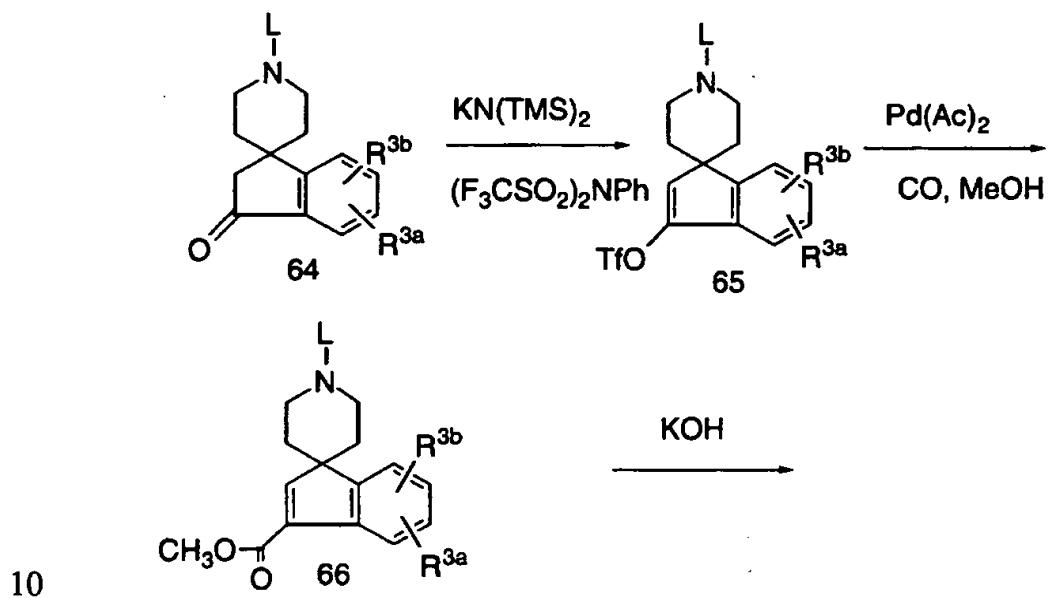
As shown in Scheme 24, when D=S, compound 57 can be oxidized to the sulfoxide 57 (n=1) and the sulfone 57 (n=2) by many oxidizing agents. For example, sodium periodate is often used for the synthesis of sulfoxides and Oxone is used for the synthesis of sulfones.

10 Removal of the protecting group provides the amine 56 which then can be incorporated into a growth hormone secretagogue via the chemistry detailed in Scheme 1 and 8 shown above which utilize generic intermediate 2.

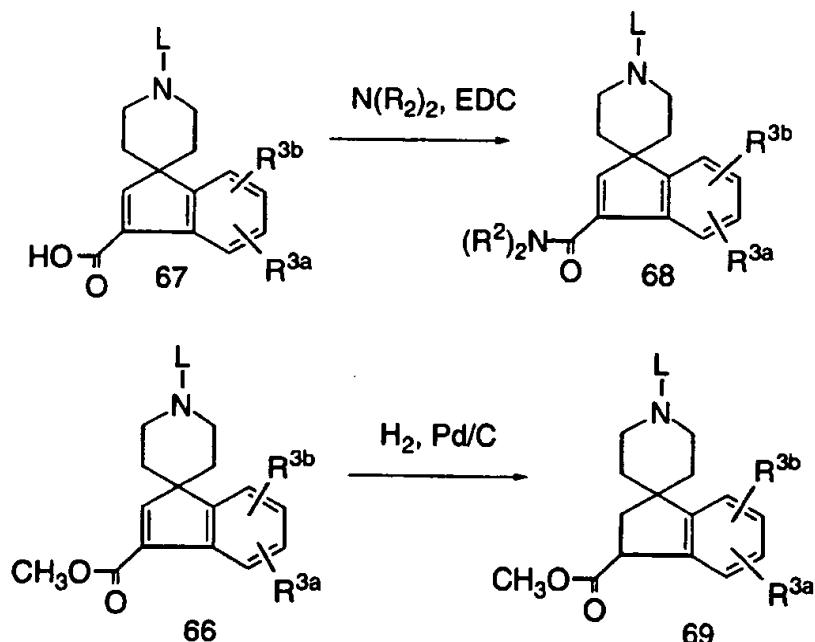
- 56 -

Compounds of formula I wherein R⁴ and R⁵ are each hydrogen can be further elaborated by reductive alkylation with an aldehyde by the aforementioned procedures or by alkylations such as by reaction with various epoxides. The products, obtained as 5 hydrochloride or trifluoroacetate salts, are conveniently purified by reverse phase high performance liquid chromatography (HPLC) or by recrystallization.

SCHEME 26



- 57 -



Homologation of the spiroindanone 64 provides easy access to spiroindanyl intermediates containing acid and ester groups. This chemistry is described in Scheme 26. Treatment of 64 with a base in an inert solvent such as THF followed by the addition of a triflating agent provides the enol triflate. Carboxylation of the enol triflate according to the procedure of Cacchi, S. *Tetrahedron Letters*, 1985, 1109-1112 provides the ester 66. The protecting group can then be removed as described above and the resulting amine can be incorporated into the subject compound via the chemistry depicted in Schemes 1 and 8. A compound containing an acid function is readily available via saponification of the ester group as the final step of the synthesis.

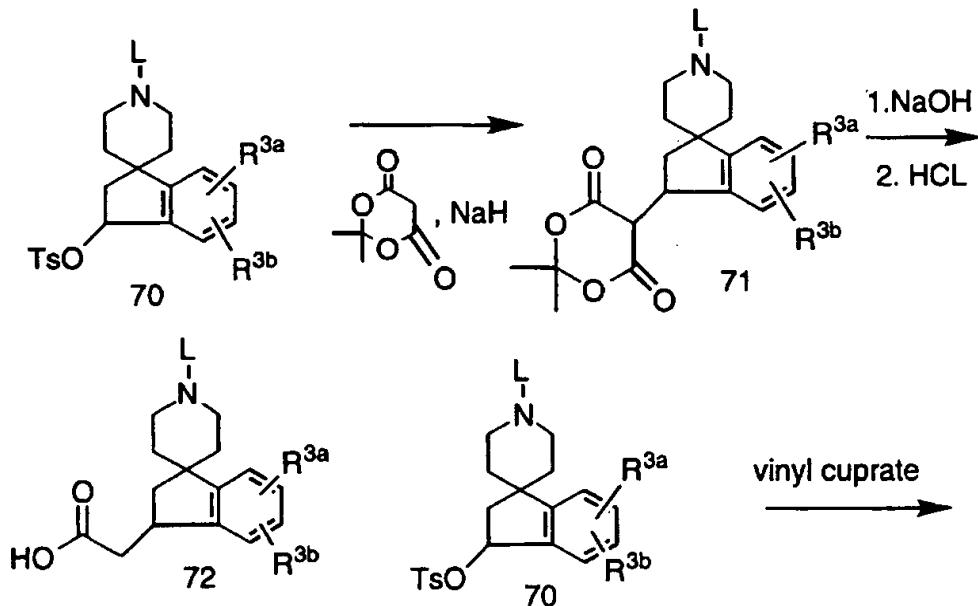
15 Saponification of the ester of 66 provides an acid which can be conveniently derivatized as for example reaction with an amine in the presence of a coupling agent such as EDC gives amides which can then be incorporated into final compounds following the chemistry detailed in Schemes 1 and 8.

Hydrogenation of 66 using a palladium catalyst in an inert solvent provides the saturated compounds which can then either be derivatized as above or carried on to the final products via the

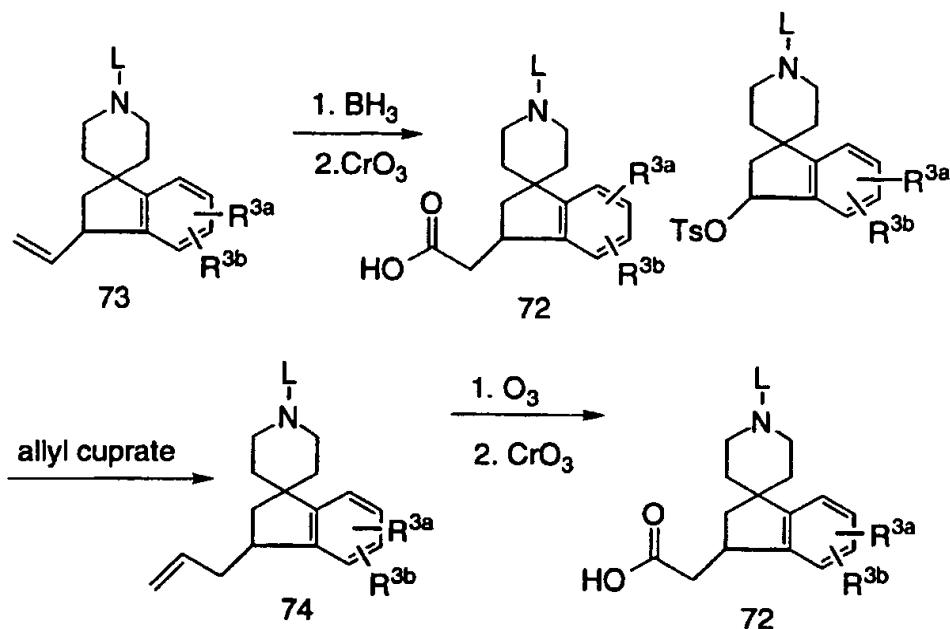
- 58 -

chemistry described in Schemes 1 and 8. The ester may also be reduced to a primary alcohol with LAH and to a aldehyde with DIBALH. Reductive alkylation of the aldehyde with ammonium acetate and sodium cyanoborohydride affords an amino methyl analog. These 5 hydroxymethyl and aminomethyl analogs may then be further reacted to afford additional growth hormone secretagogues of the general formula I. Chiral acids are available by a variety of methods known to those skilled in the art including asymmetric catalytic hydrogenation and resolution of a pair of diastereomeric salts formed by reaction with a 10 chiral amine such as D or L α -methylbenzylamine. The absolute stereochemistry can be determined in a number of ways including X-ray crystallography of a suitable crystalline derivative. 15 Spiroindane intermediates, for incorporation into growth hormone secretagogues, can be further elaborated in the benzylic position by the chemistry detailed in the following schemes.

SCHEME 27



- 59 -

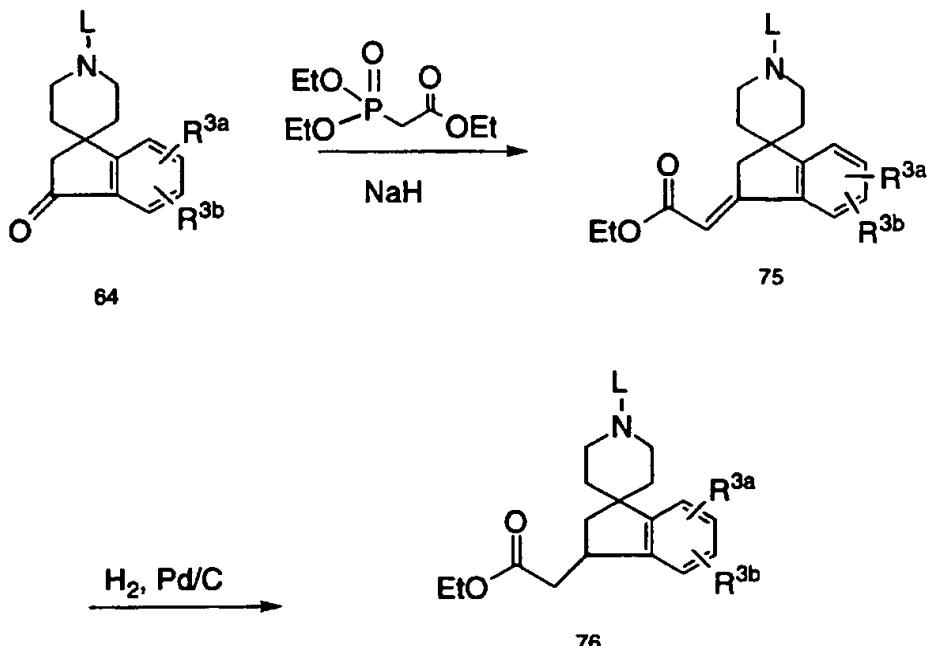


As depicted in Scheme 27, homologs of ester 69 can be conveniently prepared by a variety of methods known to those skilled in the art including the displacement of an activated alcohol such as

5 tosylate 70 by a malonate nucleophile followed by decarboxylation or a cuprate reaction followed by the adjustment of the chain length or oxidation state as appropriate.

- 60 -

SCHEME 28



Alternatively the reaction of spiroindanone 64 with Wittig or Emmons reagents also provides access to homologs of ester 69. The chemistry is described in Scheme 28. Treatment of triethylphosphonoacetate with a base in an inert solvent such as THF followed by the addition of ketone 64 provides the unsaturated ester 75. Hydrogenation of 75 using a palladium catalyst in an inert solvent provides the saturated ester 76. The protecting group can then be removed as described above and the resulting amine can be incorporated into a final compound via the chemistry described in Schemes 1 and 8. A secretagogue containing an acid function can be obtained via saponification of the ester function as the final step of the synthesis.

Chiral esters and acids are available by a variety of methods known to those skilled in the art including asymmetric catalytic hydrogenation, chromatographic resolution of a pair of diasteromers, and via crystallization of salts formed from chiral amines such as D or L- α -methylbenzylamine. The absolute stereochemistry can be

- 61 -

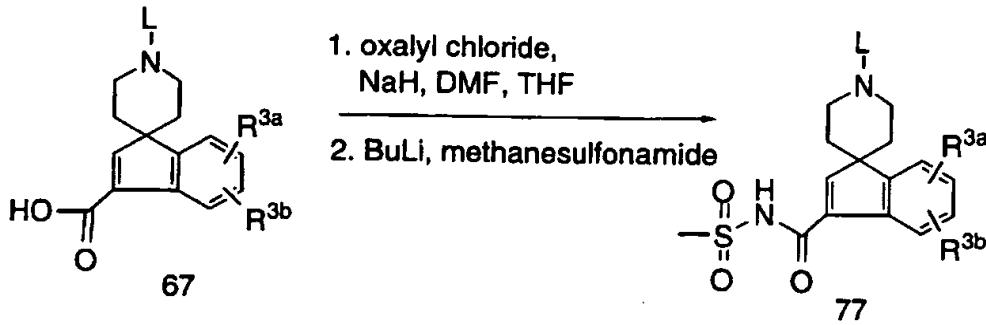
determined in a number of ways including X-ray crystallography of a suitable crystalline derivative.

The ester can be reduced to an alcohol by treatment with LAH and to an aldehyde with DIBALH. Reductive alkylation of the 5 aldehyde with ammonium acetate and sodium cyanoborohydride affords an amino methyl analog. These hydroxymethyl and aminomethyl analogs may then be further reacted to afford additional growth hormone secretagogues of the general formula 1.

Saponification of ester 44 provides an acid which can be 10 conveniently derivatized as for example reaction with an amine in the presence of a coupling reagent such as EDC gives amides which can be incorporated into a secretagogue as detailed in Schemes 1 and 8.

Homologation of ester 44 is possible using a variety of 15 methods known to those skilled in the art including the method described in *J. Org. Chem.* 1992, 57 7194-7208.

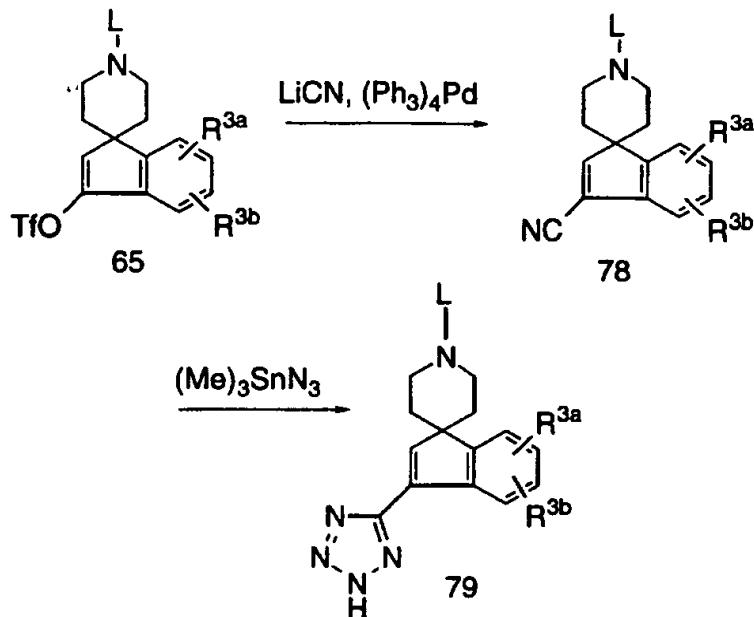
SCHEME 29



As shown in Scheme 29, a variety of acid equivalents can 20 also be incorporated into the spiroindane intermediates for example acylsulfonamides are readily available from acids such as 67 and 72. Treatment of the spiroindane acid with a base in an inert solvent such as THF followed by the addition of oxalyl chloride provides an acid 25 chloride which is then treated with a sodium salt of a sulfonamide. The protecting group can then be removed using chemistry described above and the resulting amine can be incorporated into a final compound using chemistry depicted in Schemes 1 and 8.

- 62 -

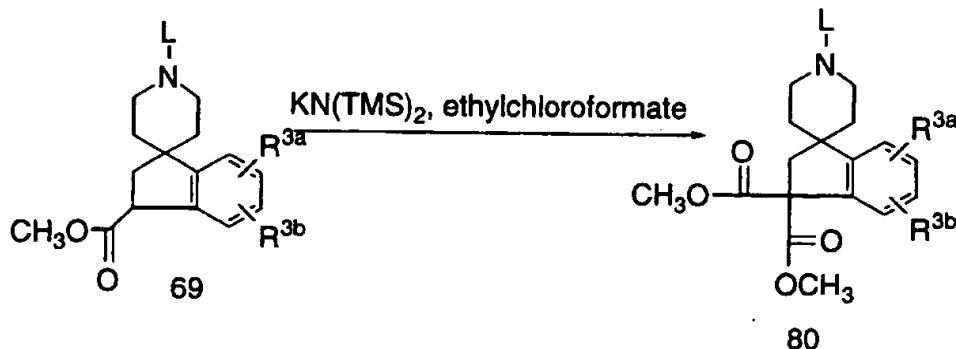
SCHEME 30



As shown in Scheme 30, tetrazole spiroindane intermediates
 5 are available from nitriles of both the shorter and longer homolog
 series. For example the reaction of enol triflate 65 with a cyanide anion
 and a palladium catalyst in the presence of an inert solvent such as
 toluene provides the unsaturated nitrile which can be converted into the
 tetrazole by reaction with trimethylstannyl azide in an inert solvent at
 10 elevated temperatures. Reduction of the indene double bond in 78 and
 79 with catalysts such as Pd/C in ethanol affords the corresponding
 saturated analogs.

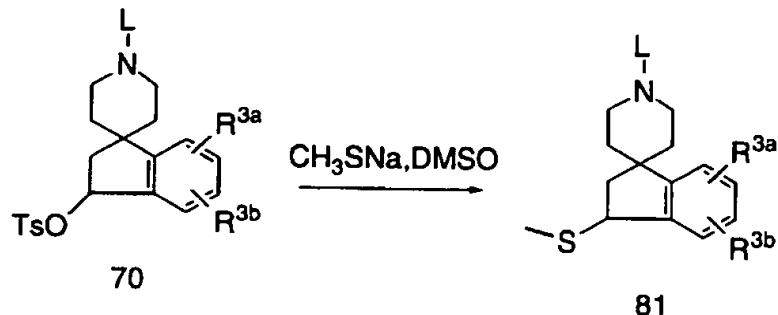
- 63 -

SCHEME 31



As shown in Scheme 31, esters such as 69 can be conveniently acylated or alkylated next to the ester function by treatment with a variety of bases and alkylating or acylating agents. For example reaction of 69 with potassium bis(trimethyl-silyl)amide in an inert solvent such as THF followed by the addition of ethyl chloroformate provides 80 in good yield. Removal of the protecting group and incorporation into the subject compounds can be accomplished as described above.

SCHEME 32

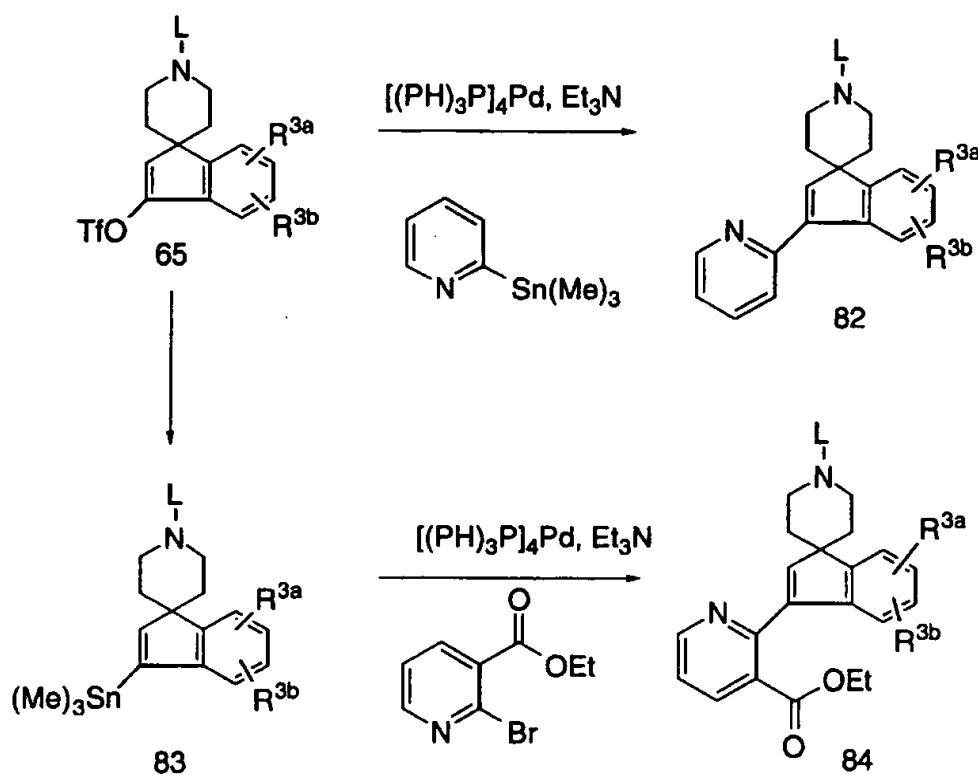


As shown in Scheme 32, further substitution at the benzylic position of the spiroindanes is readily carried out via the tosylate of the alcohol. Displacement of the tosylate with a variety of nucleophiles is possible. For example treatment of tosylate 70 with sodium thiomethoxide in DMSO provides the sulfide 81. The protecting group

- 64 -

can be removed as above and the resulting amine can be incorporated into the final compound employing chemistry described in Schemes 1 and 8. Alternatively the sulfide can be oxidized to the sulfoxide or sulfone by treatment with the appropriate oxidizing agent prior to deprotection or as the final step in the synthesis.

SCHEME 33



10 As shown in Scheme 33, the incorporation of aryl and heteroaryl groups into the benzylic position of spiroindanes is most conveniently carried out via the enol triflate 65. Palladium catalysed reaction of the enol triflate with a variety of aryl or heteroarylstannanes in an inert solvent such as toluene provides the desired intermediates.

15 For example 2-trimethylstannyl-pyridine reacts with 65 in the presence of a catalytic amount of tetrakis(triphenylphosphine)palladium in toluene at reflux to give the coupled product 82. Alternatively the enol

- 65 -

triflate 65 can be converted into the vinyl stannane 83 by reaction with hexamethylditin and a palladium catalyst in an inert solvent such as toluene. The vinyl stannane can then be coupled with a variety of aryl or hetero aryl bromides or triflates, for example coupling to 2-bromo-5 3-carbo-methoxypyridine provides 84. The protecting group L can be removed from the coupled products using chemistry described above and the resulting amine can be included in the final compound as described in Schemes 1 and 8.

10 In some cases the order of carrying out the foregoing reaction schemes may be varied to facilitate the reaction or to avoid unwanted reaction products.

15 The utility of the compounds of the present invention as growth hormone secretagogues may be demonstrated by methodology known in the art, such as an assay disclosed by Smith , et al., Science, 260, 1640-1643 (1993) (see text of Figure 2 therein). In particular, all of the compounds prepared in the following examples had activity as growth hormone secretagogues in the aforementioned assay. Such a result is indicative of the intrinsic activity of the present compounds as growth hormone secretagogues.

20 The growth hormone releasing compounds of Formula I are useful *in vitro* as unique tools for understanding how growth hormone secretion is regulated at the pituitary level. This includes use in the evaluation of many factors thought or known to influence growth hormone secretion such as age, sex, nutritional factors, glucose, amino acids, fatty acids, as well as fasting and non-fasting states. In addition, the compounds of this invention can be used in the evaluation of how other hormones modify growth hormone releasing activity. For example, it has already been established that somatostatin inhibits growth hormone release. Other hormones that are important and in need of study as to their effect on growth hormone release include the gonadal hormones, e.g., testosterone, estradiol, and progesterone; the adrenal hormones, e.g., cortisol and other corticoids, epinephrine and norepinephrine; the pancreatic and gastrointestinal hormones, e.g., insulin, glucagon, gastrin, secretin; the vasoactive peptides, e.g.,

5 bombesin, the neuropeptides; and the thyroid hormones, e.g., thyroxine and triiodothyronine. The compounds of Formula I can also be employed to investigate the possible negative or positive feedback effects of some of the pituitary hormones, e.g., growth hormone and endorphin
10 peptides, on the pituitary to modify growth hormone release. Of particular scientific importance is the use of these compounds to elucidate the subcellular mechanisms mediating the release of growth hormone.

10 The compounds of Formula I can be administered to animals, including man, to release growth hormone *in vivo*. For example, the compounds can be administered to commercially important animals such as swine, cattle, sheep and the like to accelerate and increase their rate and extent of growth, to improve feed efficiency and to increase milk production in such animals. In addition, these
15 compounds can be administered to humans *in vivo* as a diagnostic tool to directly determine whether the pituitary is capable of releasing growth hormone. For example, the compounds of Formula I can be administered *in vivo* to children. Serum samples taken before and after such administration can be assayed for growth hormone. Comparison
20 of the amounts of growth hormone in each of these samples would be a means for directly determining the ability of the patient's pituitary to release growth hormone.

25 Accordingly, the present invention includes within its scope pharmaceutical compositions comprising, as an active ingredient, at least one of the compounds of Formula I in association with a pharmaceutical carrier or diluent. Optionally, the active ingredient of the pharmaceutical compositions can comprise an anabolic agent in addition to at least one of the compounds of Formula I or another composition which exhibits a different activity, e.g., an antibiotic growth retardant or an
30 agent to treat osteoporosis or in combination with a corticosteroid to minimize the latter's catabolic side effects or with other pharmaceutically active materials wherein the combination enhances efficacy and minimizes side effects.

- 67 -

Growth promoting and anabolic agents include, but are not limited to, TRH, diethylstilbestrol, amino acids, estrogens, β -agonists, theophylline, anabolic steroids, enkephalins, E series prostaglandins, retinoic acid, compounds disclosed in U.S. Patent No. 3,239,345, e.g., 5 zeranol, and compounds disclosed in U.S. Patent No. 4,036,979, e.g., sulbenox, or peptides disclosed in U.S. Patent No. 4,411,890.

A still further use of the compounds of this invention is in combination with other growth hormone secretagogues such as the growth hormone releasing peptides GHRP-6, GHRP-1 as described in 10 U.S. Patent Nos. 4,411,890 and publications WO 89/07110, WO 89/07111 and B-HT920 as well as hexarelin and GHRP-2 as described in WO 93/04081 or growth hormone releasing hormone (GHRH, also 15 designated GRF) and its analogs or growth hormone and its analogs or somatomedins including IGF-1 and IGF-2 or α -adrenergic agonists such as clonidine or serotonin 5HT1D agonists such as sumatriptan or agents which inhibit somatostatin or its release such as physostigmine and pyridostigmine. In particular, the compounds of this invention may be used in combination with growth hormone releasing factor, an analog of 20 growth hormone releasing factor, IGF-1, or IGF-2. For example, a compound of the present invention may be used in combination with IGF-1 for the treatment or prevention of obesity. In addition, a compound of this invention may be employed in conjunction with retinoic acid to improve the condition of musculature and skin that 25 results from intrinsic aging.

25 The present invention is further directed to a method for the manufacture of a medicament for stimulating the release of growth hormone in humans and animals comprising combining a compound of the present invention with a pharmaceutical carrier or diluent.

As is well known to those skilled in the art, the known and 30 potential uses of growth hormone are varied and multitudinous. Thus, the administration of the compounds of this invention for purposes of stimulating the release of endogenous growth hormone can have the same effects or uses as growth hormone itself. These varied uses may be summarized as follows: stimulating growth hormone release in

elderly humans; treating growth hormone deficient adults; prevention of catabolic side effects of glucocorticoids; treatment of osteoporosis; stimulation of the immune system, acceleration of wound healing; accelerating bone fracture repair; treatment of growth retardation;

5 treating acute or chronic renal failure or insufficiency; treatment of physiological short stature, including growth hormone deficient children; treating short stature associated with chronic illness; treating obesity and growth retardation associated with obesity; treating growth retardation associated with Prader-Willi syndrome and Turner's

10 syndrome; accelerating the recovery and reducing hospitalization of burn patients or following major surgery such as gastrointestinal surgery; treatment of intrauterine growth retardation, and skeletal dysplasia; treatment of hypercorticism and Cushing's syndrome; treatment of peripheral neuropathies; replacement of growth hormone

15 in stressed patients; treatment of osteochondrodysplasias, Noonan's syndrome, sleep disorders, schizophrenia, depression, Alzheimer's disease, delayed wound healing, and psychosocial deprivation; treatment of pulmonary dysfunction and ventilator dependency; prevention or treatment of congestive heart failure, improving pulmonary function,

20 restoring systolic and diastolic function, increasing myocardial contractility, decreasing peripheral total vascular resistance, diminishing or preventing loss of body weight and enhancing recovery following congestive heart failure; increasing appetite; attenuation of protein catabolic response after a major operation; treating malabsorption

25 syndromes; reducing cachexia and protein loss due to chronic illness such as cancer or AIDS; accelerating weight gain and protein accretion in patients on TPN (total parenteral nutrition); treatment of hyperinsulinemia including nesidioblastosis; adjuvant treatment for ovulation induction and to prevent and treat gastric and duodenal ulcers;

30 stimulation of thymic development and prevention of the age-related decline of thymic function; adjunctive therapy for patients on chronic hemodialysis; treatment of immunosuppressed patients and to enhance antibody response following vaccination; increasing the total lymphocyte count of a human, in particular, increasing the T4/T8-cell ratio in a

- 69 -

human with a depressed T4/T8-cell ratio resulting, for example, from infection, such as bacterial or viral infection, especially infection with the human immunodeficiency virus; treatment of syndromes manifested by non-restorative sleep and musculoskeletal pain, including
5 fibromyalgia syndrome or chronic fatigue syndrome; improvement in muscle strength, mobility, maintenance of skin thickness, metabolic homeostasis, renal homeostasis in the frail elderly; stimulation of osteoblasts, bone remodelling, and cartilage growth; stimulation of the immune system in companion animals and treatment of disorders of
10 aging in companion animals; growth promotant in livestock; and stimulation of wool growth in sheep. Further, the instant compounds are useful for increasing feed efficiency, promoting growth, increasing milk production and improving the carcass quality of livestock. Likewise, the instant compounds are useful in a method of treatment of
15 diseases or conditions which are benefited by the anabolic effects of enhanced growth hormone levels that comprises the administration of an instant compound.

In particular, the instant compounds are useful in the prevention or treatment of a condition selected from the group
20 consisting of: osteoporosis; catabolic illness; immune deficiency, including that in individuals with a depressed T4/T8 cell ratio; bone fracture, including hip fracture; musculoskeletal impairment in the elderly; growth hormone deficiency in adults or in children; short stature in children; obesity; sleep disorders; cachexia and protein loss
25 due to chronic illness such as AIDS or cancer; and treating patients recovering from major surgery, wounds or burns, in a patient in need thereof.

In addition, the instant compounds may be useful in the treatment of illnesses induced or facilitated by corticotropin releasing
30 factor or stress- and anxiety-related disorders, including stress-induced depression and headache, abdominal bowel syndrome, immune suppression, HIV infections, Alzheimer's disease, gastrointestinal disease, anorexia nervosa, hemorrhagic stress, drug and alcohol withdrawal symptoms, drug addiction, and fertility problems.

- 70 -

It will be known to those skilled on the art that there are numerous compounds now being used in an effort to treat the diseases or therapeutic indications enumerated above. Combinations of these therapeutic agents some of which have also been mentioned above with 5 the growth hormone secretagogues of this invention will bring additional, complementary, and often synergistic properties to enhance the growth promotant, anabolic and desirable properties of these various therapeutic agents. In these combinations, the therapeutic agents and the growth hormone secretagogues of this invention may be independently 10 present in dose ranges from one one-hundredth to one times the dose levels which are effective when these compounds and secretagogues are used singly.

Combined therapy to inhibit bone resorption, prevent osteoporosis and enhance the healing of bone fractures can be illustrated 15 by combinations of bisphosphonates and the growth hormone secretagogues of this invention. The use of bisphosphonates for these utilities has been reviewed, for example, by Hamdy, N.A.T. Role of Bisphosphonates in Metabolic Bone Diseases. *Trends in Endocrinol. Metab.*, , 4, 19-25 (1993). Bisphosphonates with these utilities include 20 alendronate, tiludronate, dimethyl - APD, risedronate, etidronate, YM-175, clodronate, pamidronate, and BM-210995. According to their potency, oral daily dosage levels of the bisphosphonate of between 0.1 mg and 5 g and daily dosage levels of the growth hormone secretagogues of this invention of between 0.01 mg/kg to 20 mg/kg of 25 body weight are administered to patients to obtain effective treatment of osteoporosis.

In the case of alendronate daily oral dosage levels of 0.1 mg to 50 mg are combined for effective osteoporosis therapy with 0.01 mg/kg to 20 mg/kg of the growth hormone secretagogues of this 30 invention.

Osteoporosis and other bone disorders may also be treated with compounds of this invention in combination with calcitonin, estrogens, raloxifene and calcium supplements such as calcium citrate or calcium carbonate.

Anabolic effects especially in the treatment of geriatric male patients are obtained with compounds of this invention in combination with anabolic steroids such as oxymetholone, methyltestosterone, fluoxymesterone and stanozolol.

5 The compounds of this invention can be administered by oral, parenteral (e.g., intramuscular, intraperitoneal, intravenous or subcutaneous injection, or implant), nasal, vaginal, rectal, sublingual, or topical routes of administration and can be formulated in dosage forms appropriate for each route of administration.

10 Solid dosage forms for oral administration include capsules, tablets, pills, powders and granules. In such solid dosage forms, the active compound is admixed with at least one inert pharmaceutically acceptable carrier such as sucrose, lactose, or starch. Such dosage forms can also comprise, as is normal practice, additional substances other than inert diluents, e.g., lubricating agents such as magnesium stearate. In the case of capsules, tablets and pills, the dosage forms may 15 also comprise buffering agents. Tablets and pills can additionally be prepared with enteric coatings.

15 Liquid dosage forms for oral administration include pharmaceutically acceptable emulsions, solutions, suspensions, syrups, and elixirs containing inert diluents commonly used in the art, such as water. Besides such inert diluents, compositions can also include adjuvants, such as wetting agents, emulsifying and suspending agents, and sweetening, flavoring, and perfuming agents.

20 Preparations according to this invention for parenteral administration include sterile aqueous or non-aqueous solutions, suspensions, or emulsions. Examples of non-aqueous solvents or vehicles are propylene glycol, polyethylene glycol, vegetable oils, such as olive oil and corn oil, gelatin, and injectable organic esters such as ethyl oleate. Such dosage forms may also contain adjuvants such as 25 preserving, wetting, emulsifying, and dispersing agents. They may be sterilized by, for example, filtration through a bacteria-retaining filter, by incorporating sterilizing agents into the compositions, by irradiating the compositions, or by heating the compositions. They can also be

- 72 -

manufactured in the form of sterile solid compositions which can be dissolved in sterile water, or some other sterile injectable medium immediately before use.

5 Compositions for rectal or vaginal administration are preferably suppositories which may contain, in addition to the active substance, excipients such as cocoa butter or a suppository wax.

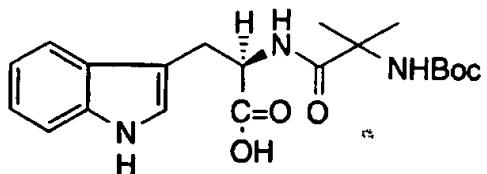
Compositions for nasal or sublingual administration are also prepared with standard excipients well known in the art.

10 The dosage of active ingredient in the compositions of this invention may be varied; however, it is necessary that the amount of the active ingredient be such that a suitable dosage form is obtained. The selected dosage depends upon the desired therapeutic effect, on the route of administration, and on the duration of the treatment. Generally, dosage levels of between 0.0001 to 10 mg/kg. of body weight daily are 15 administered to patients and animals, e.g., mammals, to obtain effective release of growth hormone. Preferably, the dosage level will be about 0.001 to about 25 mg/kg per day; more preferably about 0.01 to about 10 mg/kg per day.

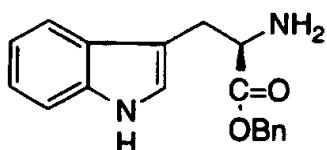
20 The following examples are provided for the purpose of further illustration only and are not intended to be limitations on the disclosed invention.

- 73 -

INTERMEDIATE 1



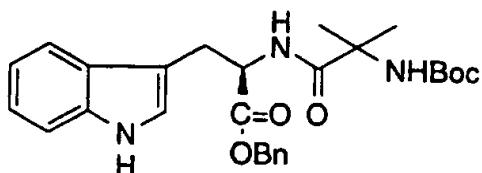
Step A:



5 To a solution of the commercially available N-t-BOC-D-tryptophan (25.0 g, 82.2 mmol), benzyl alcohol (10.2 mL, 98.6 mmol), and DMAP (100 mg) in dichloromethane (200 mL) at 0°C, was added EDC (17.4 g, 90.4 mmol) in several portions over a one hour period. The reaction mixture was stirred at room temperature for six hours and 10 was poured into water (200 mL), and the organic layer was separated. The organic solution was washed with a mixture of brine and 3 N hydrochloric acid, dried over anhydrous magnesium sulfate, filtered and concentrated to give a thick oil, which solidified upon standing.

15 To a solution of this oil in 30 mL of dichloromethane was added 20 mL of TFA and stirred for 1h. The reaction mixture was concentrated, neutralized carefully with saturated aqueous sodium bicarbonate solution, and extracted with dichloromethane (2X100 mL). The combined organic solution was washed with brine (100 mL), passed through a short column of silica gel eluting with 5-10% methanol in dichloromethane to give 23.2 g of the amine as an oil after evaporation.

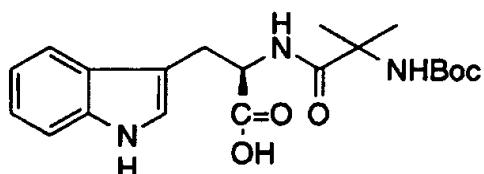
Step B:



- 74 -

To a solution of the above product, HOBT (10.6 g, 78.8 mmol) and N-BOC- α -methyl alanine (19g, 94.5 mmol) in 200 mL of dichloromethane, was added EDC (19.5 g, 0.102 mol) in several portions at 0°C. After 5 minutes, the clear reaction mixture became milky. After stirring at room temperature overnight, the reaction mixture was poured into 200 mL of water and the organic layer was separated. The organic solution was washed with brine, and with a brine and saturated sodium bicarbonate solution, dried over anhydrous magnesium sulfate, filtered and concentrated to give a thick oil, which was purified by flash chromatography eluting with 10-40% ethyl acetate in hexane to give the desired material (28.7 g). 1 H NMR (CDCl₃, 200 MHz) δ 8.48 (br.s, 1H), 7.54 (br.d, 1H), 7.38-7.23 (m, 3H), 7.19 (br.d, 2H), 7.15-7.00 (m, 1H), 6.90 (d, 1H), 6.86 (d, 1H), 5.06 (br.s, 2H), 4.95 (ddd, 1H), 3.30 (2dd, 2H), 1.40 (s, 15H)

Step C:

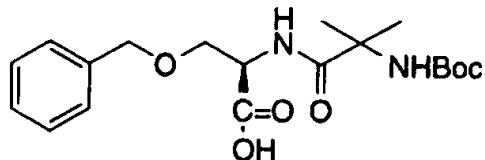


20 A solution of the material from Step B (28.7g) in 200 mL of ethanol was stirred at RT under a H₂ balloon for 20 minutes in the presence of 10% palladium on carbon (2 g). The catalyst was filtered off through a pad of celite and washed with ethyl acetate. The filtrate was concentrated to give the acid as a slightly pink foam (23.3 g).

25 ¹H NMR (CD₃OD, 400 MHz) δ 7.56 (d, J=8 Hz, 1 H), 7.31 (dd, J=1, 8 Hz, 1 H), 7.09 (s, 1 H), 7.07 (dt, J=1, 7 Hz, 1 H), 6.98 (dt, J=1, 7 Hz, 1 H), 4.69 (t, J=6 Hz, 1 H), 3.34-3.23 (m, 2 H), 1.35 (s, 3 H), 1.34 (s, 9 H), 1.29 (s, 3 H). FAB-MS calc. for C₂₀H₂₇N₃O₅ : 389; Found 390 (M+H), 290 (M+H-100 (BOC)).

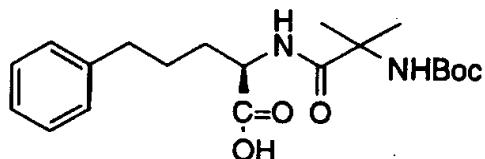
INTERMEDIATE 2

- 75 -



Following the procedures for the preparation of Intermediate 1 using N-t-Boc-O-Benzyl-D-serine in the place of N-t-BOC-D-tryptophan gave Intermediate 2. FAB-MS calc. for 5 C₁₉H₂₈N₂O₆ : 380; Found 381 (M+H), 325 (M+H-56 (t-Bu)), 281 (M+H-100 (BOC)).

INTERMEDIATE 3



10

Step A: (DL)-N-Acetyl-2-amino-5-phenylpentanoic acid

To a solution of sodium (2.3 g, 0.1 mol) in ethanol (60 mL) under nitrogen at room temperature, was added diethyl acetamidomalonate. The mixture was stirred at room temperature for one hour, and then 1-bromo-3-phenylpropane was added dropwisely. After the addition, the mixture was stirred at room temperature for two hours, then refluxed overnight. It was cooled to room temperature and partitioned between water and ethyl acetate. The organic layer was washed with sodium bicarbonate in water, dried over MgSO₄ and evaporated to give an intermediate (32.5 g, 97%). ¹H NMR (CDCl₃, 400MHz) 7.26-7.10 (m, 5 H); 6.75 (br. s, 1 H); 4.19 (q, J=7 Hz, 4 H); 2.58 (t, J=7.9 Hz, 2 H); 2.39-2.35 (m, 2 H); 2.00 (s, 3 H); 1.43-1.39 (m, 2 H); 1.20 (t, J=7 Hz, 6 H).

The product above was suspended in 190 mL of 2.5 N 25 NaOH in water and refluxed for two hours. The mixture was cooled to 0°C, and it was carefully neutralized with 6 N HCl to pH2. The precipitate was collected using a glass sinter funnel and washed with a small amount of cold water and air dried. The solid was then suspended in 300 mL of water and refluxed for four hours. The solution was

- 76 -

cooled and acidified to pH1 and the solid was collected by filtration (15.3 g, 67%). ^1H NMR (CD₃OD, 400MHz) 7.26-7.12 (m, 5 H); 4.90-4.37 (m, 1 H); 2.65-2.60 (m, 2 H); 1.97 (s, 3 H); 1.87 -1.82 (m, 1 H); 1.73-1.65 (m, 3 H).

5

Step B: (D)-N-Acetyl-2-amino-5-phenylpentanoic acid

The racemic intermediate from the previous step (10 g, 42.5 mmol) and CoCl₃-6H₂O were dissolved in 21 ml of 2 N KOH and 200 mL of water at 40°C, and the pH of the solution was adjusted to 8 by the addition of the several drops of 2 N KOH. Then acylase I (Aspergillus sp, 0.5 u/mg, from Sigma; 0.9 g) was added with vigorous stirring. The reaction mixture was stirred for one day at 40°C and the pH was kept at 8 by the addition of a few drops of KOH. The solid which formed was filtered off. The filtrate was acidified by 3 N HCl to pH2, and was extracted with ethyl acetate (200 mLX4). The organic extracts were combined and evaporated to give a white solid (4.64 g, 46%). ^1H NMR (CD₃OD, 400MHz) 7.26-7.12 (m, 5 H); 4.90-4.37 (m, 1 H); 2.65-2.60 (m, 2 H); 1.97 (s, 3 H); 1.87 -1.82 (m, 1 H); 1.73-1.65 (m, 3 H).

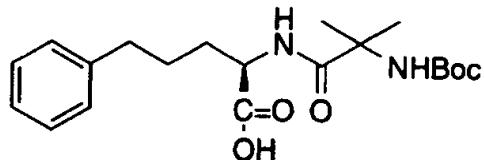
10

Step C: (D)-N-t-Boc-2-amino-5-phenylpentanoic acid

The intermediate from step B (4.2 g, 17.8 mmol) was suspended in 2 N HCl (100 mL) and refluxed for two hours. The reaction mixture was evaporated in vacuo to remove water and hydrochloric acid to yield a white solid. To a solution of this solid in 50 mL of water, was added 3 N NaOH until the pH 11, then di-t-butyl dicarbonate (4.66 g, 21.4 mmol) was added with vigorous stirring. After four hours, the reaction mixture was acidified to pH2 with 3 N HCl and it was extracted with ethyl acetate (100 mLX3). The organic extracts were combined and evaporated to give a white solid (6.56 g, crude) which was used without purification. ^1H NMR (CD₃OD, 400MHz) 7.26-7.12 (m, 5 H); 4.11-4.08 (m, 1 H); 2.65-2.60 (m, 2 H); 1.83-1.62 (m, 4 H); 1.43 (s, 9 H).

- 77 -

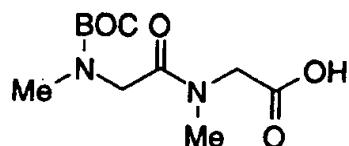
Step D:



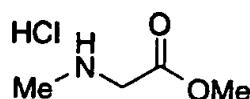
Following the procedures for the preparation of Intermediate 1 using (D)-N-t-Boc-2-amino-5-phenylpentanoic acid in the place of N-t-BOC-D-tryptophan gave Intermediate 3. ^1H NMR (CDCl_3 , 400MHz) 7.24-7.20 (m, 2H), 7.15-7.04 (m, 3H), 4.60-4.55 (m, 1H), 2.62-2.55 (m, 2H), 2.00-1.86 (m, 1H), 1.78-1.60 (m, 3H), 1.50 (s, 6H), 1.30 (s, 9H).

10

INTERMEDIATE 4



Step A:

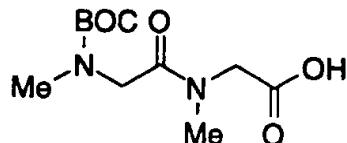


15

A solution of N-Boc Sarcosine (1.66 g) in methanol was cooled to 0°C and thionyl chloride (0.67 ml) was added slowly. After stirring for 12 hours at room temperature the mixture was concentrated to give the desired product.

20

Step B:



To a solution of the above intermediate in chloroform was added N-Boc Sarcosine (2.0 g), EDC (3.35 g), HOBr (1.42 g) and triethylamine (3.7 ml) and stirred for 12 hours at room temperature.

5 The mixture was poured into 1N HCl and extracted with methylene chloride, dried over sodium sulfate, filtered and concentrated to give 3 g of coupled product. To this material in methanol was added 6N NaOH at room temperature and stirred for 5 hours. The mixture was concentrated to remove the methanol and the residue was extracted with 10 methylene chloride. The organic layer was discarded. The aqueous layer was cooled to 0°C and acidified with conc. HCl to pH=1.0 and extracted with ethyl acetate. The organic layer was dried over sodium sulfate and concentrated to give desired product (2.0 g).

15

EXAMPLE 1

N-acetyl-*Threo*-(2R,3S)- β -methyltryptophan R-(+)- α -methylbenzyl amine salt

20 Racemic β -methyltryptophan was prepared by the method of Snyder and Matteson (*J. Am. Chem. Soc.* 1957, 79, 2217). Isomer A (100g) was suspended in 1.25L of 90/10 acetone water at 20°C and 50 mL of R-(+)- α -methylbenzylamine was added in one portion. The suspension cleared briefly before a thick white suspension formed which quickly turned to a solid mass. After aging overnight, an additional 500 25 mL of acetone was added to facilitate agitation and filtration. The suspension was filtered and the cake washed with 500 mL of acetone and sucked to a damp cake. The solid was suspended in 2.5 L of 90/10 acetone /water and heated to boiling on a steam bath. The white slurry was allowed to cool to 20°C overnight. The product was collected by 30 filtration, washed with acetone and dried yielding 39.1 g of the title compound. $\alpha = + 9.1^\circ$ (c=1, MeOH). Stereochemical assignments were made by comparison to published compounds: *J. Org. Chem.* 1994, 59, 4239 and *J. Org. Chem.* 1995, 60, 4978.

- 79 -

EXAMPLE 2

N-acetyl-*Threo*-(2S,3R)- β -methyltryptophan S-(-)- α -methylbenzyl amine salt

5 The mother liquors from Example 1 were combined and concentrated to *ca.* 1 L and 400 mL of 1 N HCl was added. The resulting suspension was stirred for 1 hr initially at 20°C then at 0°C. The product was filtered and washed with water until the filtrate was neutral. The product was sucked to a damp cake weighing 79 g. The
10 solid was suspended in 1L of 95% acetone/water and 40 mL of S-(-)- α -methylbenzylamine was added followed by 1 L of 90% acetone/water. After a few minutes a solid mass formed. An additional 500 mL of acetone was added and the mixture heated on a steam bath for *ca.* 0.5 hr. This was then allowed to stand at 20°C overnight. The product was
15 collected by filtration, washed with 500 mL of acetone, and sucked to a damp cake. The product was suspended in 2 L of 95% acetone/water and heated on a steam bath to boiling. The white suspension was allowed to cool to 20°C overnight. The product was collected by filtration, washed with 500 mL of acetone and dried yielding 54 g.
20 α = - 9.0° (c=1, MeOH).

EXAMPLE 3

N-acetyl-*Erythro* (2R,3R)- β -methyltryptophan R-(+)- α -methylbenzyl amine salt

25 170g of Isomer B (prepared as described by the reference in Example 1) which was a brittle foam containing ethyl acetate was dissolved in 2.5 L of ethyl acetate containing 100 mL of ethanol. To this was added 60 mL of R-(+)- α -methylbenzylamine. After 10 min, an additional 2L of ethyl acetate was added and the resulting thick suspension was aged at 20°C for 3 days. The product was collected by filtration, washed with ethyl acetate and and sucked to a damp cake. The salt was re-slurried four times with hot ethyl acetate containing 2%

- 80 -

water (1 x 2.5 L, 2 x 6 L, and 1 x 8 L). The yield of dried product was 43.2 g of salt. $\alpha = -19.6^\circ$ (c=1, MeOH).

EXAMPLE 4

5

N-acetyl-*Erythro* (2S,3S)- β -methyltryptophan S-(-)- α -methylbenzyl amine salt

The mother liquors from Example 3 were combined and concentrated to *ca.* 2 L and washed twice with 500 mL 1 N HCl. The 10 washes were back extracted once with ethyl acetate, and the combined ethyl acetate extracts washed twice with brine. The solution was diluted to 6 L with ethyl acetate and 60 mL of S-(-)- α -methylbenzylamine was added. After 10 min the resulting suspension was heated to boiling. The suspension was allowed to cool to ambient temperature with stirring 15 overnight. The product was collected by filtration washed with ethyl acetate and sucked to a damp cake. The salt was suspended in 6 L of ethyl acetate and suspension was heated to boiling. The suspension was allowed to cool to ambient temperature with stirring overnight. The product was collected by filtration washed with ethyl acetate and dried. 20 The yield of dried product was 65.8 g of salt. $\alpha = +19.7^\circ$ (c=1, MeOH).

EXAMPLE 5

25 N-acetyl-*threo*-(2S,3R)- β -methyltryptophan

The salt from Example 2 (53 g) was stirred with 400 mL 1 N HCl at 20°C for 20 min. The suspension was filtered and the cake washed with water until the filtrate was neutral. The wet cake was used directly for the next reaction. A sample was dried affording the title 30 compound. $\alpha = -26.4^\circ$ (c=1, MeOH).

EXAMPLE 6

threo-(2S,3R)- β -methyltryptophan

- 81 -

The wet cake from Example 5 was suspended in with 400 mL of 1 N HCl and refluxed for 12 hours. The solution was cooled to 20°C, and half of the solution was used for example 7. The title compound was isolated by adjusting the pH to 7.0 with sodium hydroxide, cooling the resulting suspension to 0°C, filtering, washing the cake with water and drying. $\alpha = -29.3^\circ$ (c=0.9, H₂O).

EXAMPLE 7

10 N-t-BOC-threo-(2S,3R)-β-methyltryptophan

The pH of the aqueous solution from Example 6 was adjusted to 7 with sodium hydroxide and cooled to 0°C. 20 g of potassium carbonate, 19 g of di-t-butyl dicarbonate, and 150 mL of THF were added. The mixture was allowed to warm slowly to ambient 15 temperature overnight. The reaction was extracted twice with ether, the aqueous acidified with 2 N HCl and extracted twice with ethyl acetate. The combined ethyl acetate extracts were washed with brine, dried with MgSO₄, filtered and concentrated affording 21.2 g of the title compound.

20

EXAMPLE 8

N-acetyl-threo-(2R,3S)-β-methyltryptophan

The title compound was prepared essentially following the 25 procedure of Example 5. $\alpha = +26.6^\circ$ (c=1, MeOH).

- 82 -

EXAMPLE 9

threo-(2*R*,3*S*)- β -methyltryptophan

5 The title compound was prepared essentially following the procedure of Example 6. $\alpha = +30.6^\circ$ ($c=0.9$, H_2O).

EXAMPLE 10

N-t-BOC-*threo*-(2*R*,3*R*)- β -methyltryptophan

10 The title compound was prepared essentially following the procedure of Example 7.

EXAMPLE 11

15 *N*-acetyl-*Erythro* (2*S*,3*S*)- β -methyltryptophan

The salt from Example 4 (65 g) was stirred with 250 mL 1 N HCl and 1.5 L of ethyl acetate at ambient temperature for 5 min. The layers were partitioned and the ethyl acetate layer was washed with 1N HCl, H_2O and brine, dried with $MgSO_4$, filtered and concentrated to afford the title compound as a brittle foam.

EXAMPLE 12

Erythro (2*S*,3*S*)- β -methyltryptophan

25 The product from Example 11 was suspended in with 500 mL of 2 N HCl and refluxed for 4 hours. The solution was cooled to 20°C, and half of the solution was used for Example 13. The title compound isolated as a foam by concentrating the solution in vacuo.

30 EXAMPLE 13

N-t-BOC-*Erythro* (2*S*,3*S*)- β -methyltryptophan

The pH of the aqueous solution from Example 6 was adjusted to 7 with sodium hydroxide and cooled to 0°C. 24 g of

- 83 -

potassium carbonate, 22 g of di-t-butyl dicarbonate, and 150 mL of THF were added. The mixture was allowed to warm slowly to ambient temperature overnight. The reaction was extracted twice with ether. The aqueous acidified with 2 N HCl and extracted twice with ethyl acetate.

5 The combined ethyl acetate extracts were washed with brine, dried with MgSO₄ filtered and concentrated. The solid was redissolved in ether, and the ether removed *in vacuo* while flushing with hexanes. The resulting slurry was filtered and dried affording 20.1 g of the title compound.

10

EXAMPLE 14

N-acetyl-*threo*-(2R,3R)- β -methyltryptophan

15 The title compound was prepared essentially following the procedure of Example 11. $\alpha = 0$ (c=1, MeOH).

EXAMPLE 15

threo-(2R,3R)- β -methyltryptophan

20 The title compound was prepared essentially following the procedure of Example 12. $\alpha = 0$ (c=0.9, H₂O).

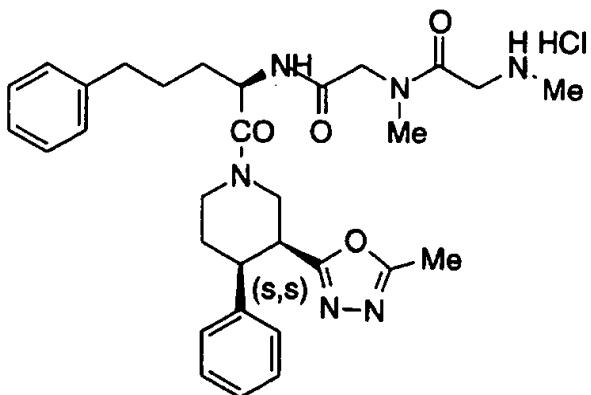
EXAMPLE 16

N-t-BOC-*threo*-(2R,3R)- β -methyltryptophan

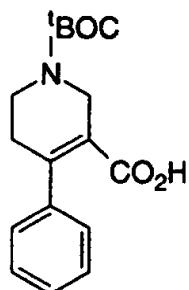
The title compound was prepared essentially following the procedure of Example 13.

- 84 -

EXAMPLE 17



Step A:



5 To a solution of 3-ethoxycarbonyl-4-piperidone hydrochloride (25 g, 120 mmole) and 3-methoxycarbonyl-4-piperidone hydrochloride (25.3 g, 130 mmole) in 70 ml of 6N aqueous sodium hydroxide was added di-t-butyl-dicarbonate (55 g, 255 mmole) in 200 ml of dioxane at 0°C. After 1/2 hour at 0°C and 3 hours at room temperature the mixture was diluted with ethyl acetate and washed with 0.5 N hydrochloric acid and brine. The organic layer was dried over magnesium sulfate, filtered and concentrated. To the crude residue in 800 ml of methylene chloride was added diisopropylethylamine (65 ml, 375 mmole) and triflic anhydride (46 ml, 275 mmole) at -78°C. After 1/2 hour the mixture was poured into saturated sodium bicarbonate solution and extracted with methylene chloride. The organic layer was washed with 1N hydrochloride, brine and dried over magnesium sulfate. The organic layer was concentrated to give the vinyl triflate. To a

10

15

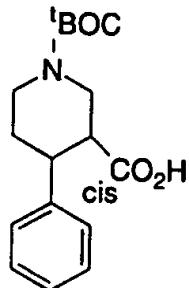
- 85 -

solution of the vinyl triflate in 700 ml of THF was added K₂CO₃ (51.7, 375 mmole), phenylboronic acid (25 g, 205 mmole) and Pd(PPh₃)₄ (3.7 g, 3.2 mmole) at room temperature. The reaction mixture was refluxed for 16 hours and then poured into water and extracted with ether (3X).

5 The organic layers were washed with water (3X), brine and dried over magnesium sulfate. The solution was then concentrated to dryness. The residue was dissolved in 100 ml of ethanol and 6 N NaOH (100 ml) was added and refluxed for 3 hours. The mixture was concentrated to dryness, the residue was dissolved in water and extracted with

10 methylene chloride. The organic layer was discarded. The aqueous layer was cooled in an ice bath and was acidified with 6 N HCl to pH=2.0. The aqueous layer was extracted with ether (3X), brine, dried over sodium sulfate and concentrated to give the desired acid (68.3 g).

15 **Step B:**

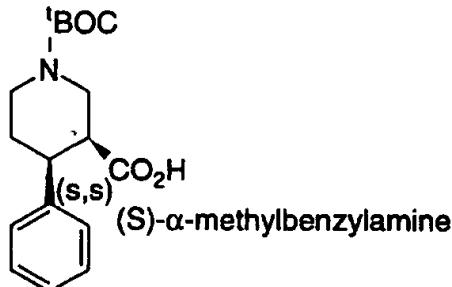


The intermediate from Step A (68.3 g, 225 mmole) was dissolved in 300 ml of methanol and hydrogenated over Pd(OH)₂ (3.7 g) at 40 psi for 6 hours and then at one atmosphere for 16 hours. The

20 mixture was filtered through Celite to remove Pd catalyst and the filtrate was concentrated under vacuum to give the cis acid (68.9 g).

- 86 -

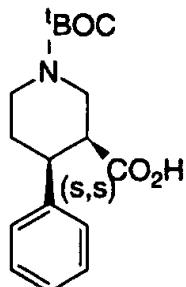
Step C:



5 The intermediate obtained in Step B (68.9 g) and (S)- α -methylbenzylamine (28.9 ml, 225 mmole) were dissolved in methanol with heating. The solution was left in refrigerator for 16 hours and the crystals that were formed were filtered off (25 g). Recrystallization from 1500 ml of ethyl acetate/methanol (5/1) gave the desired amine salt (19 g). The stereochemistry of this compound was determined by X-ray crystallographic analysis.

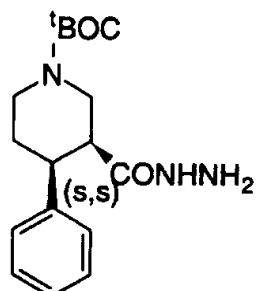
10

Step D:



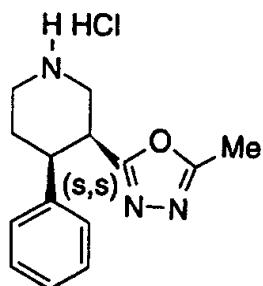
15 The intermediate from Step C (19 g) was suspended in 1N HCl and was extracted with ether. The organic extracts were combined, dried over sodium sulfate, and evaporated to give the title compound as a white solid. (13.6 g).

Step E:



To a solution of the intermediate from Step D (10 g) in THF (200 ml) was added carbonyldiimidazole (5.8 g) at room temperature. After 1 hour this mixture was slowly added to a solution of hydrazine in ethanol (200 ml) at 0°C over a 30 minutes period. The resulting mixture was warmed to room temperature and stirred an additional hour. The mixture was concentrated and the residue was taken up in water and extracted with methylene chloride, brine, dried over sodium sulfate and concentrated to give desired product (10.2 g).

Step F:

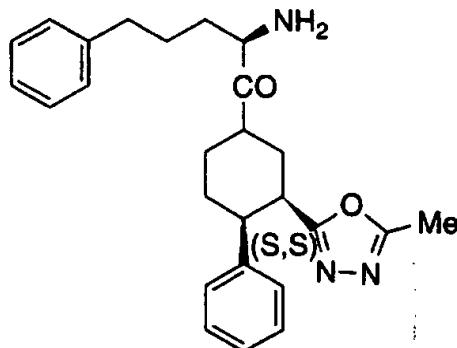


15 A solution of the intermediate from Step E (5.0g) and triethyl orthoacetate (10 ml) were heated to 100°C for 4 hours. The mixture was poured into 0.5N HCl and extracted with methylene chloride. The organic layer were washed with water, brine and dried over sodium sulfate. Concentration in vacuum and purification by 20 MPLC (hexanes/ ethyl acetate=2/1) gave the desired product which was dissolved in ethyl acetate and bubbled HCl gas at 0°C until it was

- 88 -

saturated. The mixture was stirred at room temperature for 30 minutes and evaporated to give the desired product (4.0 g).

Step G:

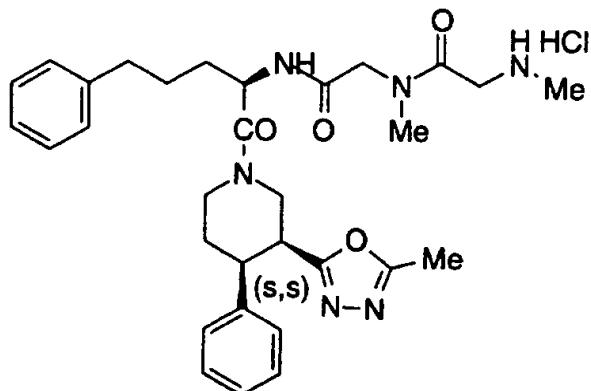


5

To a solution of the intermediate prepared in Step F (396 mg) in chloroform was added (D)-N-t-Boc-2-amino-5-phenylpentanoic acid (492 mg), EDC (541 mg), HOBr (226 mg) and triethylamine (0.58 ml) and stirred at room temperature for 12h. The mixture was poured into 1N HCl and extracted with methylene chloride, dried over sodium sulfate, filtered and concentrated. The residue was purified by RPLC (chromatotron) (hexanes/ethyl acetate=1/1) to give the coupled product (490 mg). To this coupled product in methylene chloride was added TFA and the mixture was stirred at room temperature for 1 hour. The reaction mixture was concentrated and the residue was taken up in water, basified with 1N NaOH and extracted with chloroform. The organic layer was washed with brine, dried over sodium sulfate, filtered and concentrated to give the desired product (350 mg).

- 89 -

Step H:

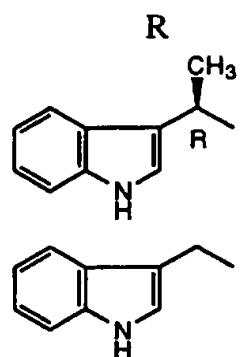
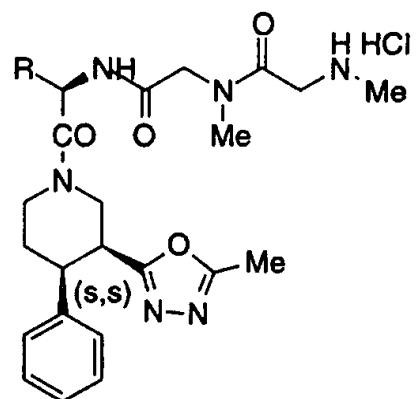


To a solution of the compound from Step G (100 mg) in chloroform was added Intermediate 4 (34 mg), EDC (53 mg) and HOBr (22 mg) and stirred at room temperature for 12h. The mixture was poured into 1N HCl and extracted with methylene chloride. The combined organics was dried over sodium sulfate, filtered and concentrated and the residue was purified by PLC (methylene chloride/methanol 20/1) to give the coupled product. To a solution of this material in ethyl acetate at 0°C was bubbled in HCl gas until it was saturated. The mixture was stirred at room temperature for 10 minutes and was evaporated to give a desired product (39 mg).

¹H NMR (400 MHz, CD₃OD, mixture of rotamers): 7.31-7.04 (m, 10 H), 4.85-4.62 (m, 2H), 4.13-3.98 (m, 5 H), 3.65 (m, 1 H), 3.37 (m, 1 H), 3.18 (m, 1 H), 3.01 (s, 3/2 H), 2.97 (s, 3/2 H), 2.74 (s, 3 H), 2.92-2.35 (m, 3 H), 2.29 (s, 3 H), 2.00-1.54 (m, 6 H). FAB-MS: 561.3 (M+1).

The following compounds were prepared according to the methods described in Steps G and H for the synthesis of Example 17 by using proper amino acids.

- 90 -

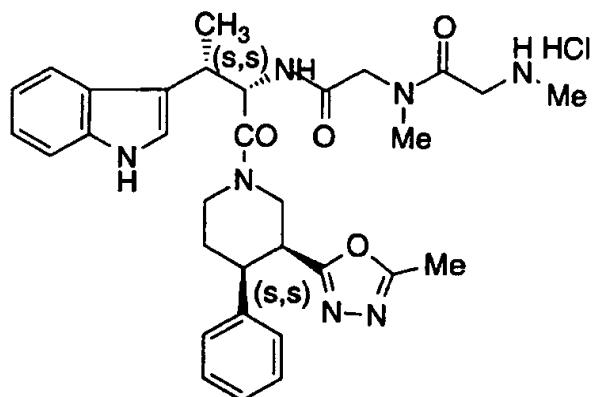


FAB-MS (M+1)
586.3

572.2

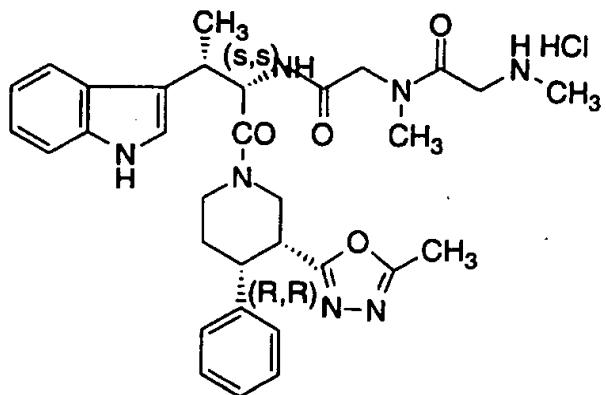
- 91 -

EXAMPLE 18



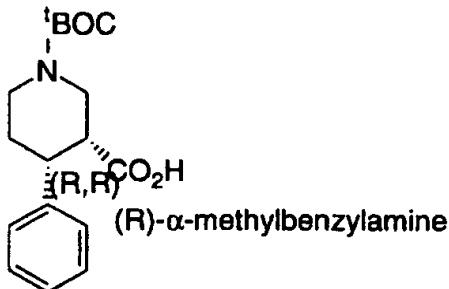
Prepared as described in Example 17 but by using N-Boc-(2S, 3S)-2-methyl-Trp instead of (D)-N-t-Boc-2-amino-5-phenylpentanoic acid. FAB-MS: 586.3 (M+1).

EXAMPLE 19



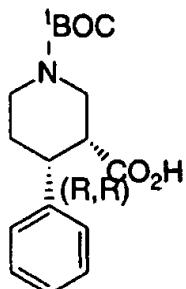
- 92 -

Step A:



The mother liquid from Step C of Example 17 was concentrated to give an off-white solid (68 g). This material was 5 dissolved in 1N NaOH and extracted with methylene chloride. The organic layer was discarded. The aqueous layer was acidified with conc. HCl at 0°C to pH=1.0 and the precipitate that had formed was collected after stirring at 0°C for 10 minutes. The white solid was dried under nitrogen to give 28 g of acid. The acid was dissolved in hot 10 methanol and (R)- α -methylbenzylamine (11.12g) was added. The solution was left in refrigerator for 16 hours and the crystals that had formed were filtered off (15 g).

Step B:

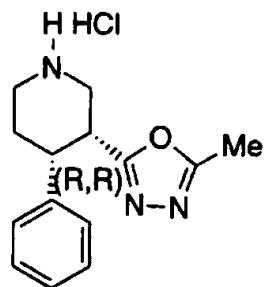


15

The intermediate from Step A (4.3 g) was suspended in 1N HCl and was extracted with ether. The extracts were combined, dried over sodium sulfate and evaporated to give a white solid. (3.0g).

- 93 -

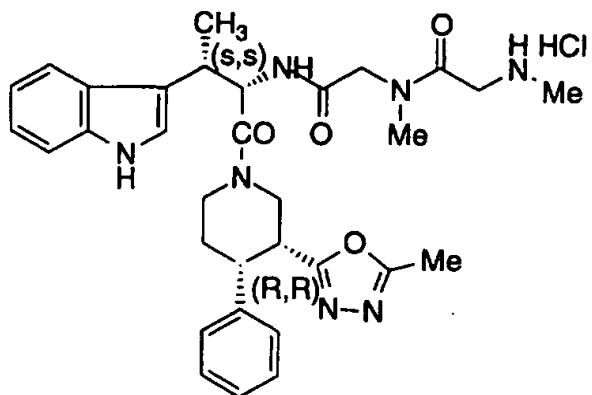
Step C:



Prepared from the intermediate (750 mg) from Step B by using the procedure described in Steps E and F of Example 17.

5

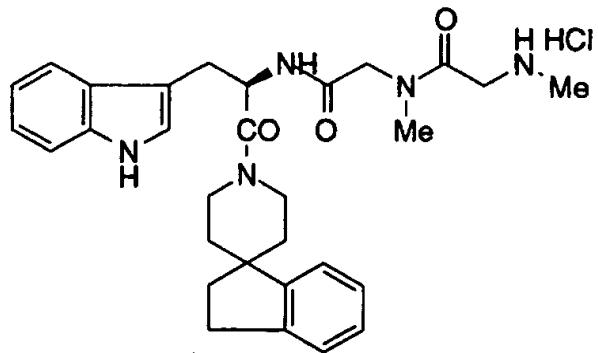
Step D:



Prepared from the intermediate synthesized in Step C by using the procedures described in Steps G and H of Example 17 (N-Boc-
10 β -(S)-methyl-(L)-Trp was used instead of (D)-N-t-Boc-2-amino-5-phenylpentanoic acid). FAB-MS: 586.3 (M+1).

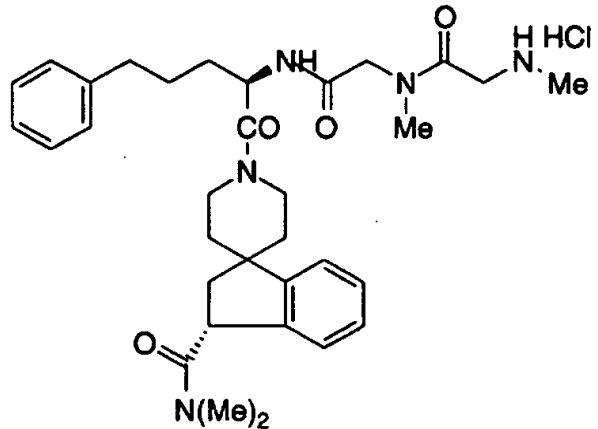
- 94 -

EXAMPLE 20



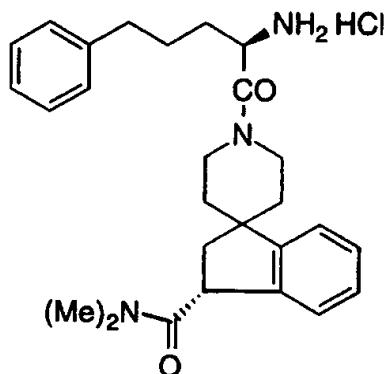
Prepared from spiro-indanyl piperidine (see *J. Med. Chem.* 1992, 35, 2033 for preparation) and N-Boc-(D)-Trp as detailed in Steps 5 G and H of Example 17. FAB-MS: 516.2 (M+1).

EXAMPLE 21



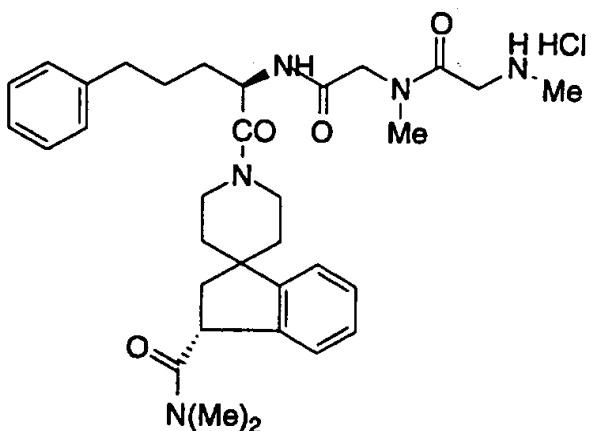
- 95 -

Step A:



The intermediate from Example 38 Step G was deprotected with ethyl acetate / HCl and the resulting product (200 mg, 0.68 mmol) 5 was coupled to (2R)-N-t-Boc-5-phenylpentanoic acid (240 mg, 0.81 mmol) using EDC (190 mg, 1mmol), HOBT (130 mg, 1.0 mmol) and NMM (0.097 ml, 0.88 mmol). The BOC protected title compound (130 mg) was obtained after purification (Prep TLC dichlormethane / acetone 9 : 1). The title compound was obtained after treatment with ethyl 10 acetate / HCl.

Step B:

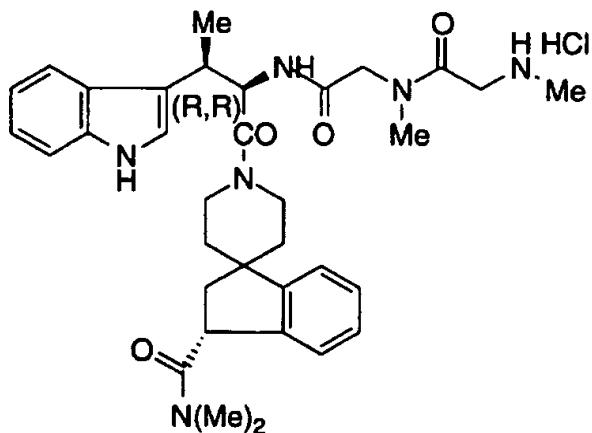


Prepared from the intermediate obtained from Step A by 15 the procedure described in Example 17, Step G and Step H to give the desired compound. ¹H NMR (400 MHz, CD₃OD, mixture of rotamers):

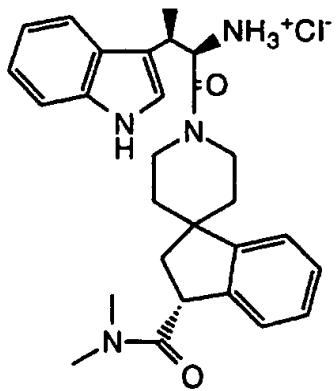
- 96 -

7.30-7.05 (m, 9 H), 4.92 (m, 1 H), 4.58 (m, 1H), 4.49 (m, 1H), 4.27-3.84 (m, 6H), 3.33 (m, 1/2 H), 3.20 (m, 1/2 H), 3.04 (s, 3 H), 3.02 (s, 3 H), 2.99 (s, 3/3 H), 2.96 (s, 3/3 H), 2.94 (s, 3/3 H), 2.74 (s, 3 H), 3.00-2.50 (m, 3 H), 2.16 (m, 1 1/2 H), 1.90-1.42 (m, 7 1/2 H). FAB-MS: 5 576.4 (M+1).

EXAMPLE 22



Step A:



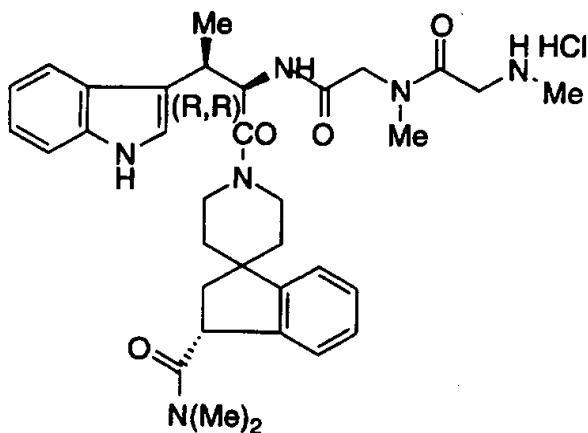
10

A sample of the title compound from Example 38 Step G was deprotected with a saturated solution of HCl in ethyl acetate to give the hydrochloride salt (6.3 g, 21 mmol). To this salt in dichloromethane at 0°C was added (R,R)-b-methyltryptophan (7.0 g, 22 mmol), HOBT 15 (4.4g, 33 mmol), NMM (4.83 ml,44 mmol) and finally EDC(6.3g, 33

- 97 -

mmol). The reaction mixture was warmed to room temperature and stirred overnight. It was then poured into ethyl acetate and washed with 1 N HCl, saturated bicarb, and brine then dried over magnesium sulfate. The organic layer was filtered and concentrated. Purification by flash chromatography (ethyl acetate) provided the title compound (10 g, 17.9 mmol). A solution of this intermediate (1.32 g, 2.6 mmol) in ethyl acetate (8 mL) was cooled to 0°C. While stirring, HCl-EtOAc was added to the mixture (10 mL). The reaction was stirred for 20 minutes, until TLC analysis indicated that the reaction was complete. The solution was then concentrated to remove the ethyl acetate to afford 1.25g of the product (100%). ESI-MS calc. for C₂₈H₃₃N₄O₂: 457; Found 458 (M+H).
10
15

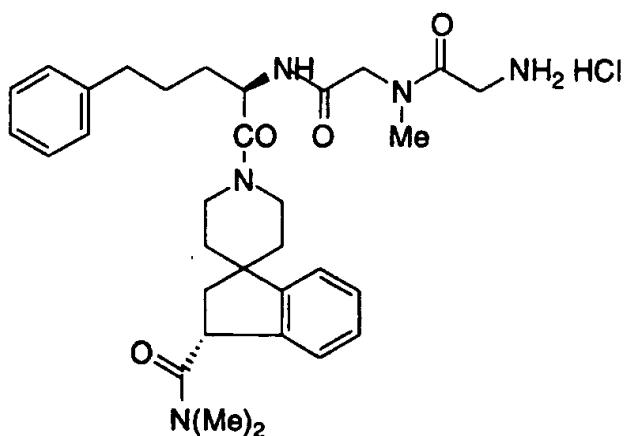
Step B:



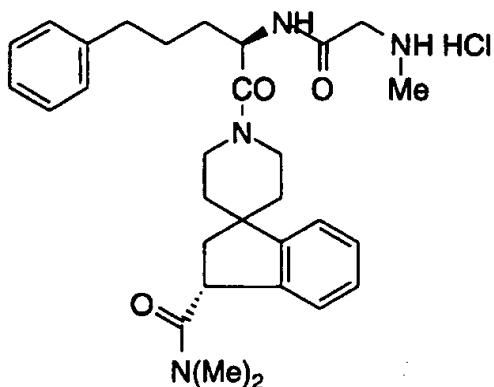
Prepared from the intermediate obtained from Step A by the procedure described in Example 17, Step G (using N-Boc-b-(R)-methyl-(D)-Trp instead of (D)-N-t-Boc-2-amino-5-phenylpentanoic acid) and Step H to give the desired compound. FAB-MS: 601.5 (M+1).

- 98 -

EXAMPLE 23



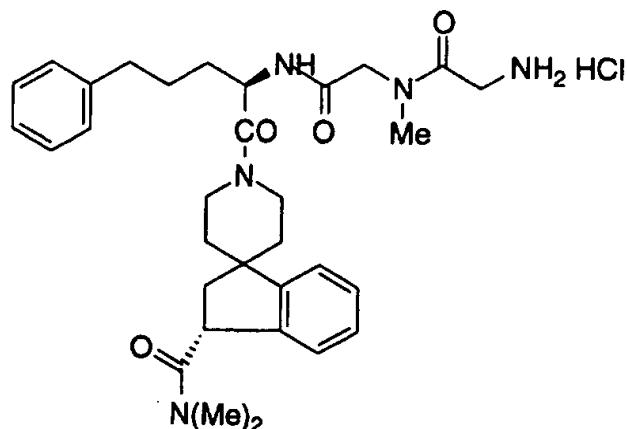
Step A:



5 Prepared from the intermediate obtained from Example 21, Step A by standard peptide-type coupling but N-Boc-Sarcosine was used instead of Intermediate 4.

- 99 -

Step B:

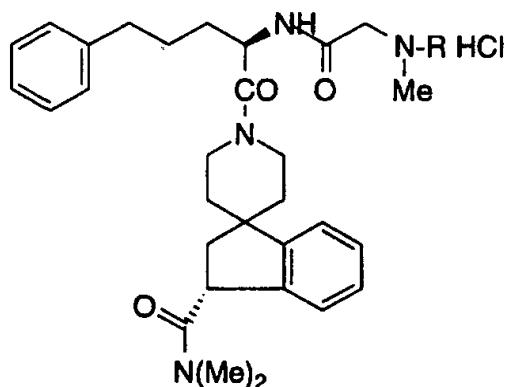


To a solution of the intermediate prepared from Step A (11 mg) in chloroform was added N-Boc-Gly (4.3 mg), EDC (8 mg), HOBT (3.3 mg) and triethylamine (0.008 ml) and stirred for 12 hours. The reaction mixture was poured into 1N HC, extracted with methylene chloride, dried over sodium sulfate, filtered and concentrated. The residue was purified by PLC (methylene chloride/methanol=10/1) to give the coupled product. This coupled product was deprotected by the HCl/ethyl acetate protocol to give a desired product (11 mg).

¹H NMR (400 MHz, CD3OD, mixture of rotamers): 7.27-7.04 (m, 9 H), 4.92 (m, 1 H), 4.56 (m, 1H), 4.49 (m, 1H), 4.24-3.87 (m, 6H), 3.33 (m, 1/2 H), 3.20 (m, 1/2 H), 3.04 (s, 3 H), 3.02 (s, 3 H), 2.99 (s, 3/2 H), 2.97 (s, 3/2 H), 3.00-2.50 (m, 3 H), 2.16 (m, 1 1/2 H), 1.90-1.42 (m, 7 1/2 H). FAB-MS: 562.4 (M+1).

- 100 -

Additional compounds were prepared as described in Example 23 Step B using the appropriate amino acids.

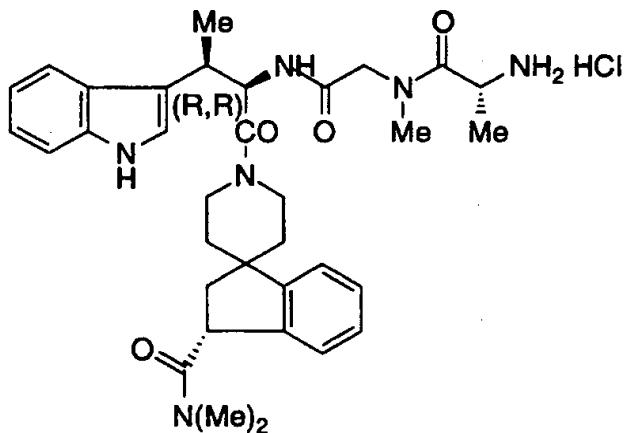


R
D-Ala
L-Ala

FAB-MS:
576.4
576.4

5

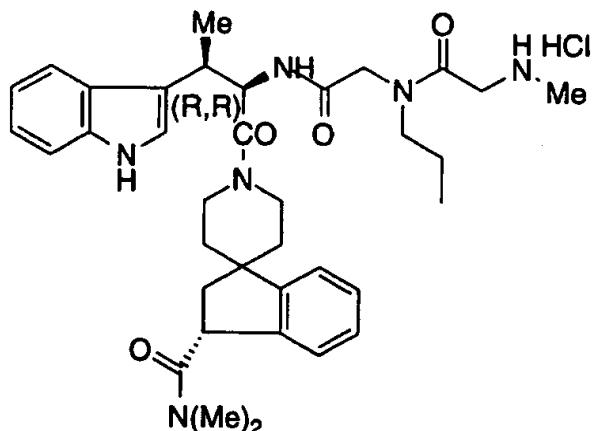
EXAMPLE 24



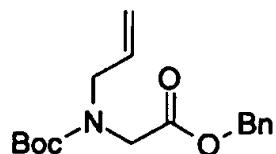
Prepared from the intermediate obtained from Example 22, Step A by using peptide-type coupling methods as described above.
10 FAB-MS: 601.6 (M+1).

- 101 -

EXAMPLE 25



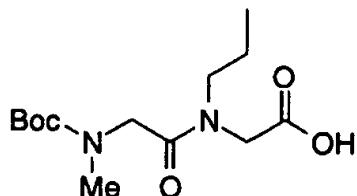
Step A:



5 To a solution of N-Boc-Gly (3.0 g) in methylene chloride (100 ml) was added benzyl alcohol (1.6 ml), EDC (4.4 g) and a catalytic amount of DMAP. After 2 hours, the mixture was poured into 1N HCl and extracted with methylene chloride, dried over sodium sulfate, filtered and concentrated to give N-Boc-Gly-OBn (4.5 g). To a solution of N-Boc-Gly-OBn (1.32 g) in THF (10 ml) was added potassium bis(trimethylsilyl)amide (1.19 g) in THF (20 ml) at -78°C. After 30 minutes at -78°C allyl bromide (0.52 ml) was added to the reaction mixture slowly. The resulting mixture was slowly warmed up to room temperature and stirred for an additional 12 hours. The mixture was 10 poured into 1N HCl and extracted with ether, the combined organics were dried over sodium sulfate, filtered and concentrated. The residue was purified by chromatatron (hexanes/ethyl acetate=10/1) to give 15 desired product (585 mg).

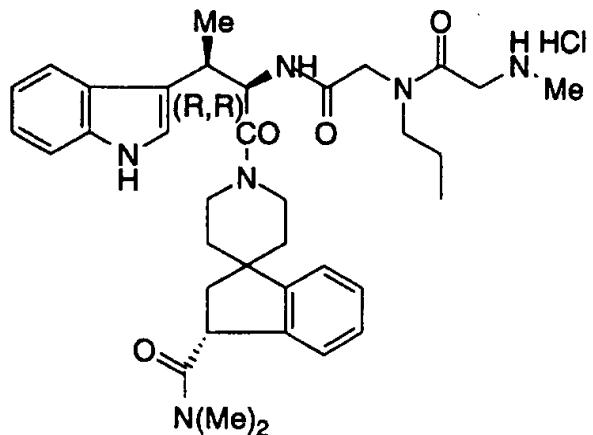
- 102 -

Step B:



The intermediate obtained from Step A (585 mg) was treated TFA with at room temperature for 10 minutes. The mixture 5 was concentrated and azeotroped with toluene (3X) to give a white solid. To this white solid in chloroform was added EDC (729 mg), N-Boc-Sar (435 mg), triethylamine (0.78 ml) and a catalytic amount of DMAP. After 12 hours the mixture was poured into 1N HCl and extracted with 10 methylene chloride, the combined organics were dried over sodium sulfate, filtered and concentrated. The residue was purified by PLC (hexanes/ethyl acetate=5/1) to give the coupled product (115 mg). This material (115 mg) in methanol was hydrogenated over Pd(OH)2 at 1 ATM for an hour. The mixture was filtered through Celite to remove 15 Pd catalyst and the filtrate was concentrated under vacuum to give the desired product (88 mg).

Step C:

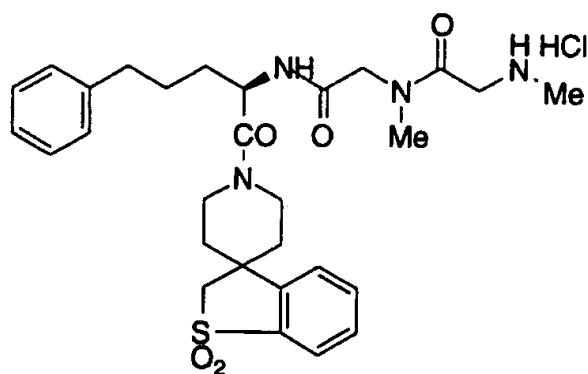


- 103 -

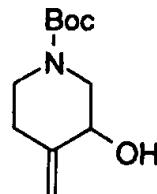
Prepared from the intermediate obtained from Example 22, Step A and the compound synthesized in Step B by standard coupling and deprotection chemistry. FAB-MS: 629.3 (M+1).

5

EXAMPLE 26



Step A:



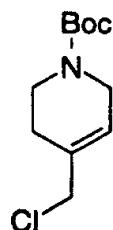
To a suspension solution of methyltriphenyl-phosphonium iodide (30 g, 74 mmole) in 150 mL of THF was slowly added butyllithium (2.5 N, 25.5 mL, 63.7 mmole) at 0°C. After an hour at room temperature, N-t-BOC protected 4-piperidone (prepared from 4-piperidone monohydrate hydrochloride by the procedure described in Protective Groups in Organic Synthesis T. W. Greene, John Wiley and Sons, NY. 1981.) in 50 mL of THF was added to reaction mixture at room temperature slowly. This reaction was stirred for 2 hours and filtered. The filtrate was concentrated and purified (MPLC, silica gel, hexanes/ethyl acetate=10/1) to give the Wittig product (7.9 g) in 82% yield. To a suspension of selenium dioxide/silica gel (prepared according to the procedure described in *Chem. Lett.* 1981, 1703) in 30 mL methylene chloride was added t-butyl hydroperoxide (1.23 mL).

- 104 -

After 15 minutes, the Wittig product (0.72 g, 3.69 mmole) in 5 mL of methylene chloride was added. The cloudy solution was stirred for 3 hours and filtered though Celite. The filtrate was washed with water, brine and dried over sodium sulfate. The organic layer was

5 concentrated and purified by flash chromatography (hexanes/ethyl acetate=4/1) to give the desired compound in 52% yield (0.41 g).

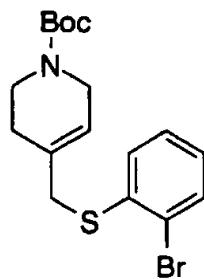
Step B:



10 The intermediate obtained from Step A (400 mg, 1.88 mmole) was dissolved in 10 mL benzene and thionyl chloride (0.165 ml, 2.26 mmole) was added and heated to 60°C for 25 minutes. The resulting mixture was poured into NaHCO₃ (aq.) and extracted with ether. The ether layer was dried over magnesium sulfate and

15 concentrated to give desired compound (333 mg, 77%).

Step C:



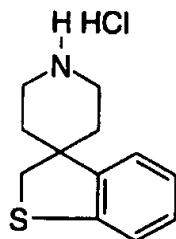
20 The intermediate obtained from Step B (330 mg, 1.43 mmole) was dissolved in 10 mL of acetone and 2-bromothiophenol (0.172 ml, 1.43 mmole) and potassium carbonate (390 mg, 2.86 mmole) were added. The reaction mixture was heated to 60°C for an hour and

- 105 -

then filtered though silica gel (100% ether). The organic layer was concentrated and purified by flash chromatography (silica gel, hexanes/ethyl acetate=10/1) to give the desired compound in 84% yield (460 mg).

5

Step D:

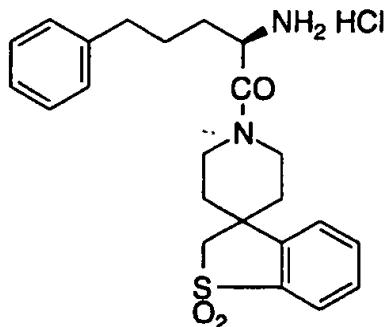


10 The intermediate obtained from Step C (450 mg, 1.17 mmole) was dissolved in 60 mL of benzene and AIBN (10 mg) and tributyltin hydride (0.644 mL, 2.39 mmole) were added. This mixture was refluxed for 2 hours and concentrated. The residue was dissolved in ether and bromine was added till the reaction solution turned to a brownish color. To this brownish solution at room temperature was added DBU (0.65 mL) in dropwise manner. The resulting cloudy 15 solution was filtered though silica gel and washed with ether. The ether solution was concentrated and the residue was purified by radial chromatography (silica gel, hexanes/ethyl acetate=10/1) to give desired compound (157 mg) in 43% yield. ^1H NMR (400 MHz, CDCl_3): 7.18 (d, 7 Hz, 1 H), 7.12 (t, 7 Hz, 1 H), 7.06 (m, 2 H), 4.11 (m, 2 H), 3.30 (s, 20 3 H), 2.89 (m, 2 H), 1.79 (m, 4 H), 1.47 (s, 9 H).

25 The cyclized product was dissolved in ethyl acetate and HCl gas was bubbled at 0°C until it was saturated. The mixture was stirred at room temperature for 30 minutes and evaporated to give the desired product.

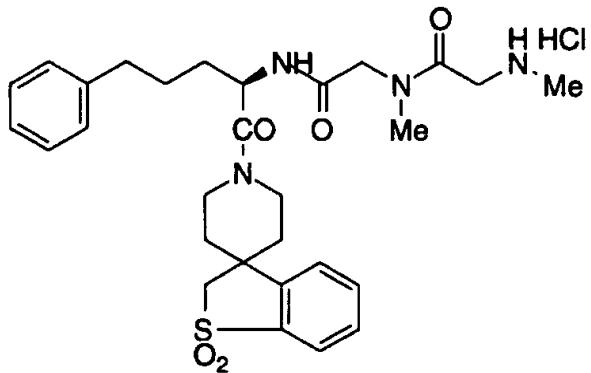
- 106 -

Step E:



To a solution of the intermediate prepared in Step D (724 mg) in chloroform was added (D)-N-t-Boc-2-amino-5-phenylpentanoic acid (873 mg), EDC (1.14 g), HOBr (445 mg) and diisopropylethylamine (0.57 ml). After 12 hours at room temperature the mixture was poured into 1N HCl and extracted with methylene chloride. The combined organics were dried over sodium sulfate, filtered and concentrated. The residue was purified by chromatatron (hexanes/ethyl acetate=1/1) to give the coupled product. To this product in methanol was added OXONE (4.6 g) at 0°C and stirred for 1 hour. The mixture was poured into water and extracted with ether. The ether extracts were dried over sodium sulfate, filtered and concentrated to give sulfone. The N-Boc group was removed by treatment with HCl gas at 0°C in ethyl acetate(1.03 g).

Step F:

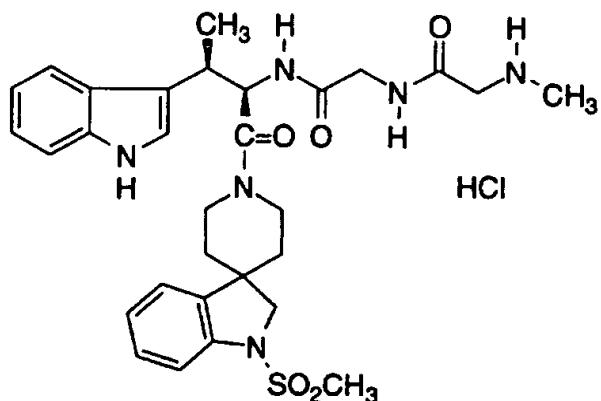


- 107 -

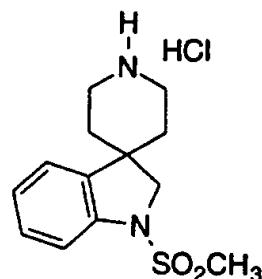
The title compound was prepared from the intermediate obtained from Step E (20 mg) by standard peptide-type coupling and deprotection chemistry. FAB-MS: 555.0 (M+1).

5

EXAMPLE 27



Step A:



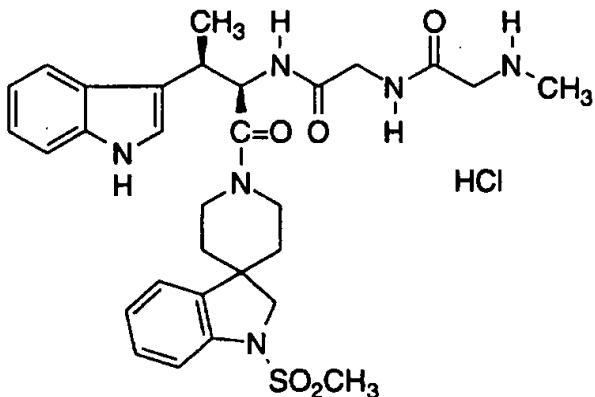
To a solution of 1.20 g (5.8mmol) of 1'-methyl-1,2-dihydro-spiro[3H-indole-3,4'-piperidine] (prepared as described in H. Ong *et al* *J. Med. Chem.* 1983, 23, 981-986) in 20 mL of dry dichloromethane at 0°C was added triethylamine (0.90 mL; 6.4 mmol) and methanesulfonyl chloride (0.49 mL; 6.35 mmol) and stirred for 30 min. The reaction mixture was poured into 15 mL of saturated aqueous sodium bicarbonate solution and extracted with dichloromethane (2X10 mL). The combined organics were washed with brine (20 mL), dried over anhydrous potassium carbonate, filtered and the solvent removed

- 108 -

under reduced pressure to yield 1.44 g of the methanesulfonamide derivative as pale yellow oil which was used without purification.

To a solution of above crude product in 20 mL of dry 1,2-dichloroethane at 0°C was added 1.0 mL (9.30 mmol) of 1-chloroethyl chloroformate, and then stirred at RT for 30 min and finally at reflux for 1h. The reaction mixture was concentrated to approximately one third of the volume and then diluted with 20 mL of dry methanol and refluxed for 1.5h. The reaction was cooled to RT and concentrated to approximately one half of the volume. The precipitate was filtered and washed with a small volume of cold methanol. This yielded 1.0 g of the piperidine HCl salt as a white solid. The filtrate was concentrated and a small volume of methanol was added followed by ether. The precipitated material was once again filtered, washed with cold methanol, and dried. This gave an additional 0.49 g of the desired product. Total yield 1.49 g (70%). ^1H NMR(CDCl₃, 200MHz) δ 7.43-7.20 (m, 3H), 7.10 (dd, 1H), 3.98 (bs, 2H), 3.55-3.40 (bd, 2H), 3.35-3.10 (m, 2H), 2.99 (s, 3H), 2.15 (t, 2H), 2.00 (t, 2H).

Step B:



20

To a suspension of 1.2 g of the piperidine prepared in Step A in 15 mL of acetonitrile was added 0.50 mL of N-methylmorpholine, 1.00 g of (2R, 3R)-N-tBOC-2-methyl-tryptophan (synthesized as described in Example), 0.80 g of HOBT, and 1.00 g of EDC and stirred at RT for 3h. The reaction mixture was diluted with 100 mL of

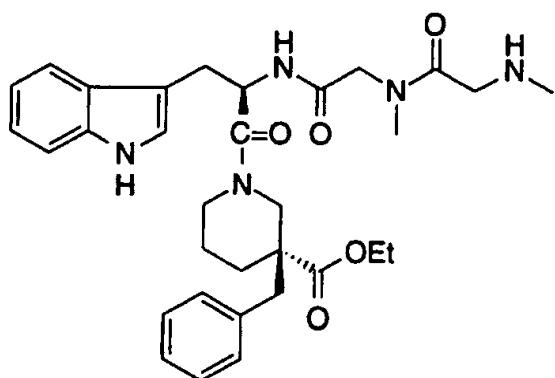
- 109 -

ether and washed with 50 mL of 0.05N HCl, 50 mL of saturated sodium bicarbonate solution, dried over MgSO₄, and concentrated. A solution of the above intermediate in 50 mL of ethyl acetate at 0°C was treated with HCl (g) for 2 min. and then stirred for 1h. Dry ether (50 mL) was 5 added, and the precipitated solid was collected by filtration. The yield was 1.44 g.

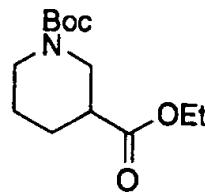
10 The above compound was coupled with Intermediate 4 under the standard peptide-type coupling reaction conditions as described above. Purification of the crude product by flash chromatography with dichloromethane-acetone 1:1 as the eluent gave the BOC compound that was deprotected by the HCl/ethyl acetate protocol to give the title compound as a pale yellow solid. ¹H NMR (CDCl₃; 400MHz).

15

EXAMPLE 28



Step A:



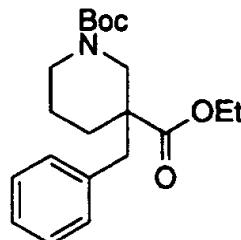
20 To a stirred solution of ethyl nipecotate (15 g, 95.4 mmol) and DMAP (0.05 eq.) in dichloromethane at 0°C was added dropwise by an addition funnel di-tert-butyl dicarbonate (21.8 g, 100 mmol) in

- 110 -

dichloromethane (200 mL). The mixture was stirred for 2-3 hours. The solution was washed with 3 N HCl and saturated sodium chloride, dried over anhydrous magnesium sulfate; then filtered and concentrated to give the desired product (18.7 g, 88%).

5

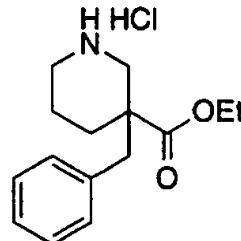
Step B:



To a stirred solution of ethyl N-t-Boc nipecotate (7 g, 26.90 mmol) in THF (100 mL) at -78°C under argon was added LHMDS (28 mL, 28 mmol) over a 10 minute period. The solution was allowed to stir an additional 30 minutes at -78°C; then benzyl bromide (4.8 g, 28 mmol) was added slowly to the solution. The reaction mixture was stirred overnight and allowed to warm to room temperature. The material was concentrated, then diluted with water, and extracted using ethyl acetate (2 x 200 mL). The organic layer was dried over anhydrous magnesium sulfate, filtered, and concentrated. Purification by silica gel flash column chromatography, eluting with 20% ethyl acetate in hexane, provided the title compound. (8.32 g, 88%). FAB-MS calc. for C₂₀H₂₉NO₄: 347; Found 348 (M+H).

10

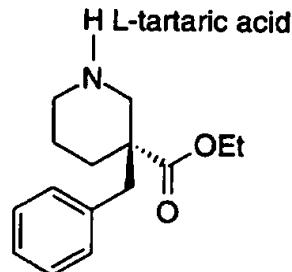
Step C:



- 111 -

A solution of the intermediate from Step B (8 g, 23.02 mmol) in ethyl acetate (80 mL) was cooled to 0°C. While stirring, hydrogen chloride gas was bubbled into the mixture until saturation occurred. The reaction was stirred for 40 minutes, until TLC analysis indicated that the reaction was complete. The solution was then concentrated to remove the ethyl acetate to afford the product (6.53 g, 99%). ¹H NMR (CDCl₃, 400MHz) δ 7.25-7.19 (m, 3 H), 7.04-7.01 (m, 2 H), 5.35 (v. br. s, 2 H), 4.22-4.10 (m, 2 H), 3.44 (d, J = 13 Hz, 1 H), 3.21 (br. d, J = 12.7 Hz, 1 H), 2.95 (d, J = 13.5 Hz, 1 H), 2.76-2.68 (m, 3 H), 2.22 (br. d, J = 13 Hz, 1 H), 1.73-1.71 (m, 1 H), 1.61-1.48 (m, 2 H), 1.18 (t, J = 7 Hz, 3 H). FAB-MS calc. for C₁₅H₂₁NO₂: 247; Found 248 (M+H).

Step D:



15

The intermediate from previous step (50.8 g) was dissolved in dichloromethane and it was washed with 3N NaOH. The aqueous layer was extracted with dichloromethane and the combined solution was dried (MgSO₄) and evaporated to give the free amine as an oil.

20 The ethyl 3-benzyl nipecotate and D-tartaric acid (31 g) were dissolved in 880 mL of water/acetone (1:4) solution with heating. The solution was left in the refrigerator overnight and the crystals which were formed were filtered off. Recrystallization in 470 mL of water/acetone (1:4) at room temperature gave the ethyl 3-(R)-benzyl nipecotate D-tartaric acid salt (21 g).

25

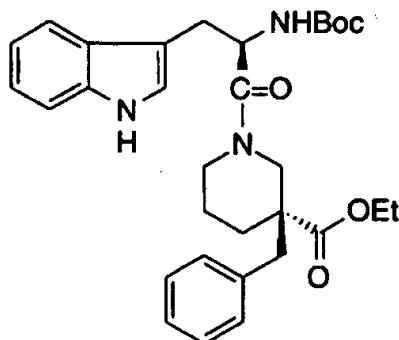
The combined mother liquor was evaporated and to it was added 3N NaOH and dichloromethane, the mixture was stirred for 30 minutes and the organic layer was separated. The aqueous was extracted

- 112 -

twice with dichloromethane and the combined organic extracts were dried over MgSO₄ and evaporated to give 24.4 g of the S-isomer enriched compound. It was crystallized with L-tartaric acid (14.8 g) in 400 mL of water/acetone (1:4) at room temperature to give ethyl 3 (S)-5 benzyl nipecotate L-tartaric acid salt (27.3 g). ¹H NMR (CD₃OD, 400MHz) δ 7.31-7.22 (m, 3 H), 7.12-7.09 (m, 2 H), 4.40 (s, 2 H, from tartaric acid), 4.30-4.10 (m, 2 H), 3.49 (br. d, J = 13 Hz, 1 H), 3.06 (d, J = 13.6 Hz, 1 H), 2.98 (d, J = 13 Hz, 1 H), 2.92 (dt, J = 3.3 Hz, 13 Hz, 1 H), 2.82 (d, J = 13.6 Hz, 1 H), 2.30 (d, J = 12.4 Hz, 1 H), 1.88 (td, J = 3 Hz, 14.5 Hz, 1 H), 1.69 (dt, J = 3 Hz, 13 Hz, 1 H), 1.63-1.51 (m, 1 H), 1.25b (q, J = 7.1 Hz, 3 H).

10

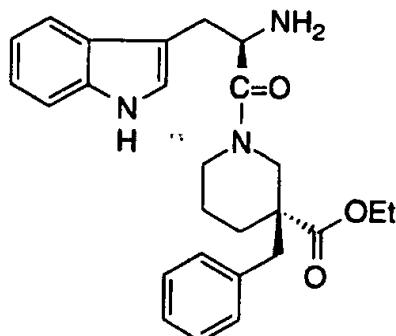
Step E:



15 Ethyl 3 (S)-benzyl nipecotate L-tartaric acid salt (39.74 g) was suspended in 70 mL of 3N NaOH and 70 mL of water, followed by extraction with dichloromethane. The extracts were combined, dried, and evaporated to give a thick oil. To a stirred solution of the oil, N-t-Boc D-TrpOH (30.43 g) and HOBr (13.5 g) in dichloromethane (200 mL) at 0°C, was added EDC (23 g) in several portions. The mixture was stirred overnight and it was poured into water and 3 N HCl and was extracted with dichloromethane. The organic layer was washed with brine, saturated sodium bicarbonate, dried over MgSO₄ and evaporated to give a crude product (67.7 g), which was used without further purification. FAB-MS calc. for C₃₁H₃₉N₃O₅: 533; Found 534 (M+H)
20
25

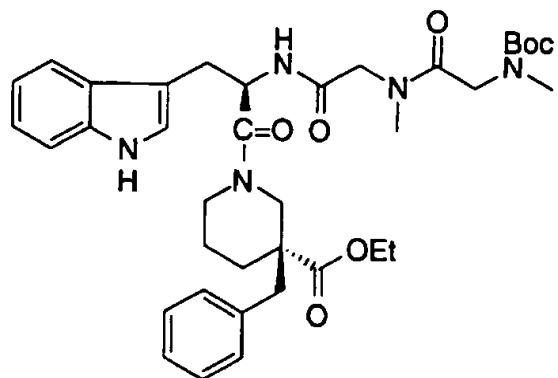
- 113 -

Step F:



To a solution of the intermediate from the previous step (67.7 g crude) in ethyl acetate (100 mL) at 0°C, was bubbled HCl gas until it was saturated. The reaction mixture was stirred at 0°C for 30 minutes and was evaporated to remove excess HCl and ethyl acetate. The residue was suspended in dichloromethane and was washed with a mixture of 3 N NaOH (70 mL) and water (100 mL). The organic layer was dried (MgSO₄), evaporated to a small volume and used in next step without further purification. FAB-MS calc. for C₂₆H₃₁N₃O₃: 433; Found 434 (M+H)

Step G:

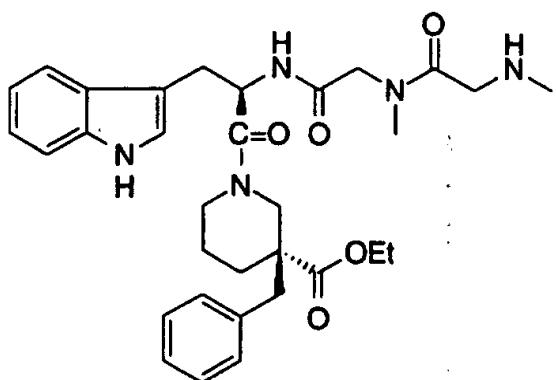


A solution containing the intermediate obtained in the last step (150 mg, 0.319 mmol), N-Boc-Sar-SarOH (83 mg, 0.319 mmol), and HOBT (43 mg) in dichloromethane (100 mL) was stirred at room

- 114 -

temperature and to it was added EDC (123 mg). The reaction mixture was stirred overnight and was worked up by diluting it with dichloromethane and washing with 3 N HCl, brine, and saturated sodium bicarbonate solution. The organic layer was dried over MgSO₄, and 5 evaporated. This residue was purified by MPLC eluting with 7% methanol in ethyl acetate to give the desired compound (188 mg, 87%)

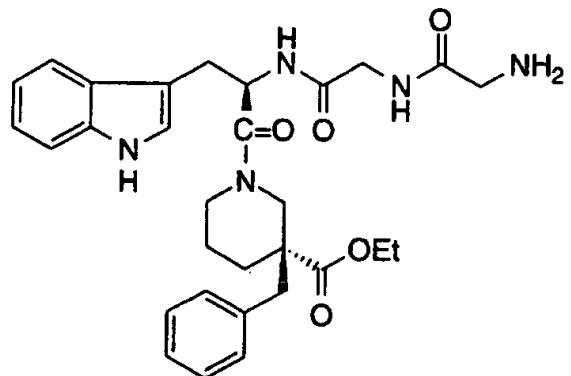
Step H:



10 To a solution of the intermediate from the previous step (160 mg, 0.237 mmol) in ethyl acetate (10 mL) at 0°C, was bubbled HCl gas until it was saturated. The reaction mixture was stirred at 0°C for 15 minutes and was evaporated to remove excess HCl and ethyl acetate to give the salts as white solid (138 mg, 95%). FAB-MS calc. for 15 C₃₂H₄₁N₅O₅: 575; Found 576 (M+H)

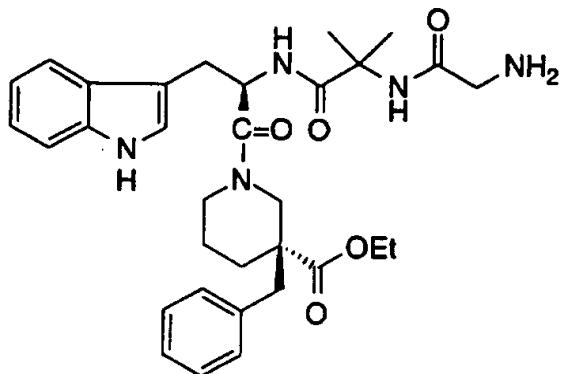
EXAMPLE 29

- 115 -

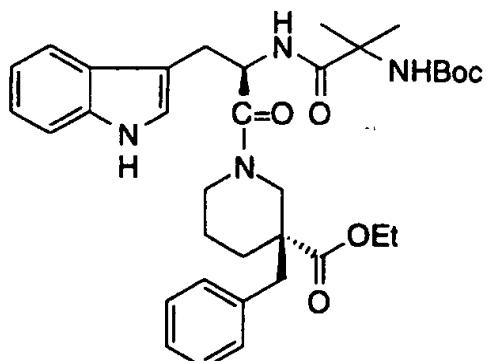


The title compound was prepared according to the similar procedures as described in Example 28, but by using Glycine instead of Sarcosine. FAB-MS calc. for C₃₀H₃₇N₅O₅: 547; Found 548 (M+H)
5

EXAMPLE 30

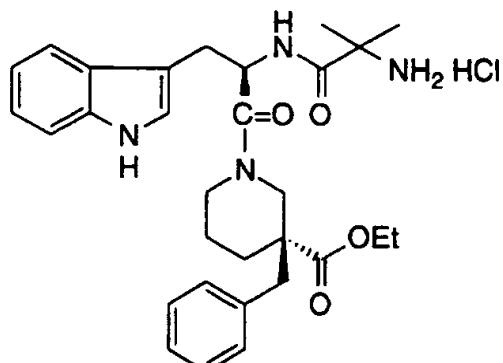


Step A:



A solution containing the intermediate obtained in the Step F, Example 28, Boc-Aib-OH (Aib: alpha-amino-isobutyric acid, 20.3 g), and DMAP (200 mg) in dichloromethane (100 mL) was stirred at room temperature and to it was added EDC (24 g) in several portions. The reaction mixture was stirred for 3 hours and was worked up by diluting it with dichloromethane and washing with 3 N HCl, brine, and saturated sodium bicarbonate solution. The organic layer was dried over MgSO₄, and evaporated to give a thick oil. This oil was passed through a pad of silica gel, eluting with 60% ethyl acetate in hexane to remove some very polar impurities, to give the desired compound (54.2 g). FAB-MS calc. for C₃₅H₄₆N₄O₆: 618; Found 619 (M+H)⁺

15 Step B:

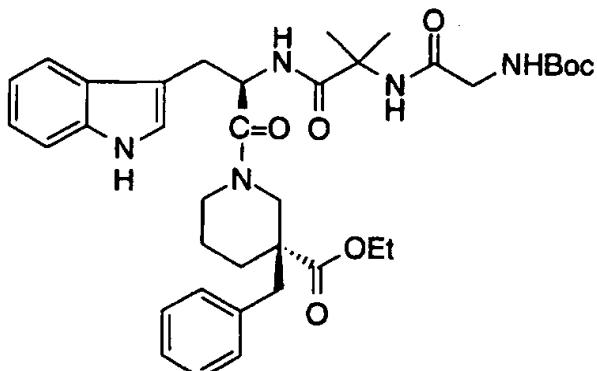


- 117 -

To a solution of the intermediate from the previous step (54.2 g) in ethyl acetate (100 mL) at 0°C, was bubbled HCl gas until it was saturated. The reaction mixture was stirred at 0°C for 15 minutes and was evaporated to remove excess HCl and ethyl acetate. The residue 5 was first dissolved in dichloromethane (100 mL) and then ethyl acetate (300 mL) was added. The mixture was stirred overnight and the solid was collected by filtration to give the title compound (34 g). Further evaporation of the mother liquor to a small volume gave the second crop of product (10.1 g). FAB-MS calc. for C₃₀H₃₈N₄O₄: 518;

10 Found 519 (M+H) ¹H NMR (CD₃OD, 400MHz) compound exists in two rotamers in approximately a 5/3 ratio that slowly interconvert relative to the NMR time scale d 7.60 (d, J = 7.9 Hz, 5/8 H), 7.55 (d, J = 7.9 Hz, 3/8 H), 7.34-6.93 (m, 9H), 5.36 (dd, J = 5.2Hz, 9.7 Hz, 3/8 H), 5.31 (dd, J = 6.7 Hz, 8.8 Hz, 5/8 H), 4.23 (br. d, J = 13.7 Hz, 3/8 H), 15 4.10-4.00 (m, 6/8 H), 4.04-3.98 (m, 3/8 H), 3.96-3.82 (m, 10/8 H), 3.80 (br. d, J = 13.5 Hz, 5/8 H), 3.36 (br. d, J = 13.3 Hz, 5/8 H), 3.29-3.22, 3.17-3.10, (2m, 2H), 3.20 (br. d, J = 14.5 Hz, 3/8 H), 3.10-2.96 (m, 5/8 H), 2.90 (s, 6/8 H), 2.60 (d, J = 13.4 Hz, 5/8), 2.41 (d, J = 13.4 Hz, 5/8 H), 2.19-2.12, 1.82-1.70, 1.68-1.60, 1.50-1.40, 1.34-1.25, 1.05-0.95 20 (6m, 4 H), 1.55 (s, 9/8 H), 1.50 (s, 15/8 H), 1.09 (t, J = 7.1 Hz, 3 H).

Step C:

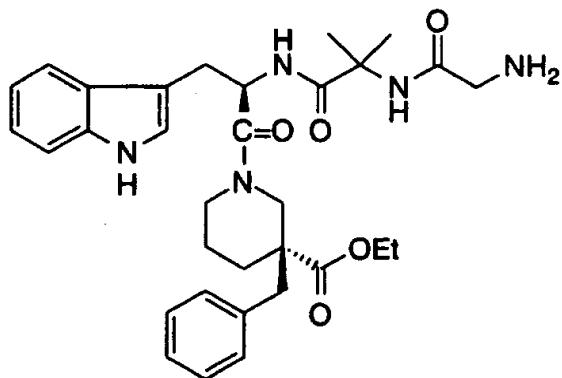


A stirred solution of the intermediate obtained in the
25 previous step (100 mg, 0.18 mmol), N-Boc-GlyOH (32 mg), and HOBT

- 118 -

(25 mg) in dichloromethane (10 mL) was added EDC (69 mg). The reaction mixture was stirred overnight and was worked up by diluting it with dichloromethane and washing with 3 N HCl, brine, and saturated sodium bicarbonate solution. The organic layer was dried over MgSO₄, 5 and evaporated. This residue was purified by MPLC eluting with ethyl acetate to give the desired compound (105 mg, 86%).

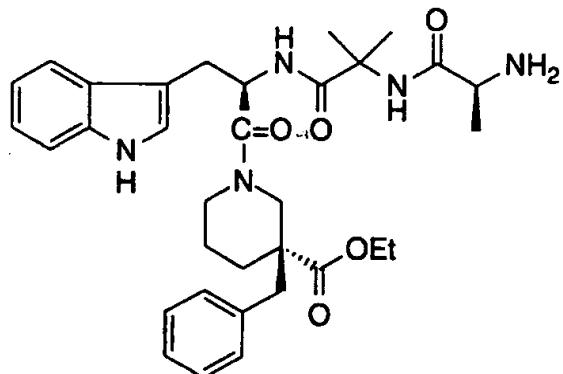
Step D:



10 To a solution of the intermediate from the previous step (105 mg, 0.155 mmol) in ethyl acetate (10 mL) at 0°C, was bubbled HCl gas until it was saturated. The reaction mixture was stirred at 0°C for 15 minutes and was evaporated to remove excess HCl and ethyl acetate to give the salts as white solid (91mg, 96%). FAB-MS calc. for 15 C₃₂H₄₁N₅O₅: 575; Found 576 (M+H).

- 119 -

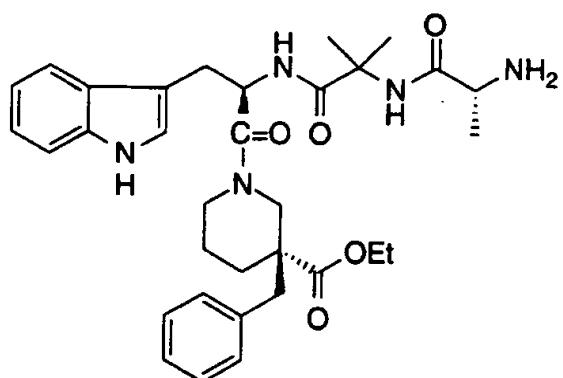
EXAMPLE 31



5 The title compound was prepared according to the same procedure as described above in Examples 28-30, but using the appropriate Boc protected amino acids which were subsequently deprotected as described above. FAB-MS calc. for C₃₃H₄₃N₅O₅: 589; Found 590 (M+H)

10

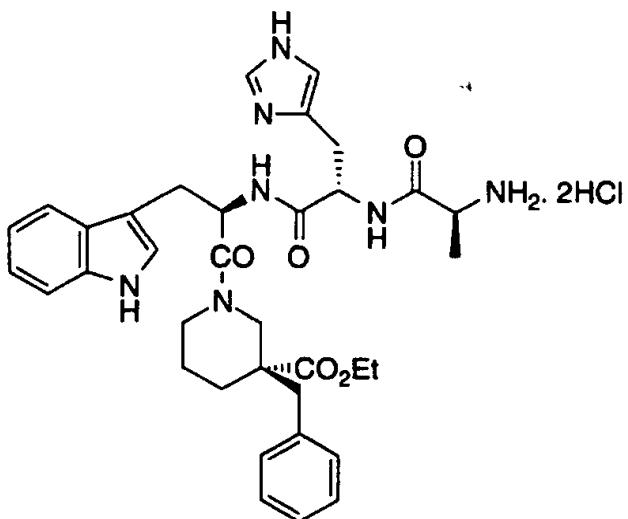
EXAMPLE 32



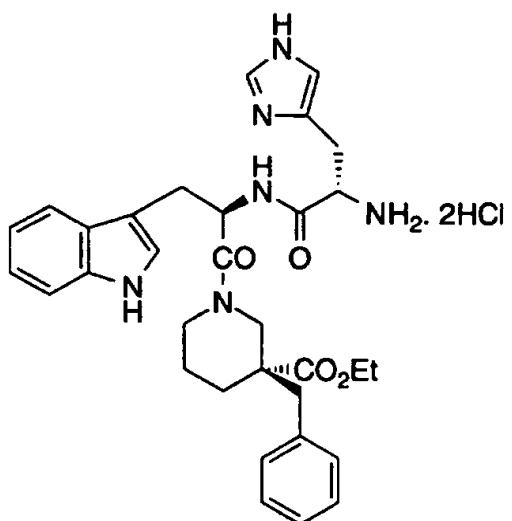
15 The title compound was prepared according to the same procedure as described above in Examples 28-30, but using the appropriate Boc protected amino acids which were subsequently deprotected as described above. FAB-MS calc. for C₃₃H₄₃N₅O₅: 589; Found 590 (M+H)

- 120 -

EXAMPLE 33



5 Step A:



To a cold solution of the product from Example 28, Step F (223 mg) in CH₂Cl₂ (3 ml) was added N-methyl-morpholine (0.075 ml), Boc-His(Boc) (205 mg) [obtained from dicyclohexylammonium salt 10 of Boc-His(Boc)], HOBT (114 mg) and EDC (134 mg; 0.699 mmol).

- 121 -

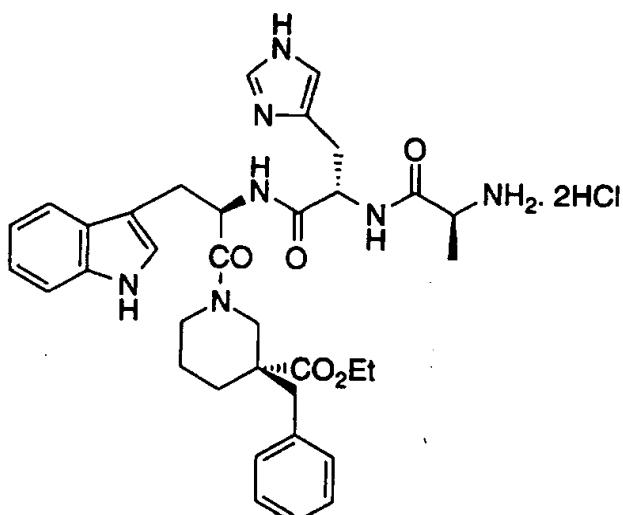
The reaction mixture was stirred at room temperature for 2 hours and then partitioned between a mixture of EtOAc/H₂O. The organic phase was separated and washed with sat. NaHCO₃, water, 10% citric acid, water, then dried over MgSO₄, filtered and concentrated in vacuo to

5 give the crude product as a foam. Purification of the crude product by flash chromatography using ethyl acetate as the eluant provided 283 mg of the protected dipeptide. The dipeptide (283 mg) was dissolved in THF (5 ml) and the solution was saturated with dry HClg for 1 minute. The reaction was sealed and stirred at room temperature for 30

10 minutes. The solvent and excess reagent were removed *in vacuo*, and the resulting residue was triturated with EtOAc to give the titled compound (240 mg).

FAB-MS: m/e calculated for C₃₂H₃₈N₆O₄: 570; found: 571 (M+H).

15 **Step B:**



To a cold (0°C) solution of the titled compound from Step A (126 mg) in CH₂Cl₂ (2 ml) were added Boc-L-Ala (47 mg), HOBT (50 mg), N-methylmorpholine (49 ml) and EDC (58 mg). The reaction

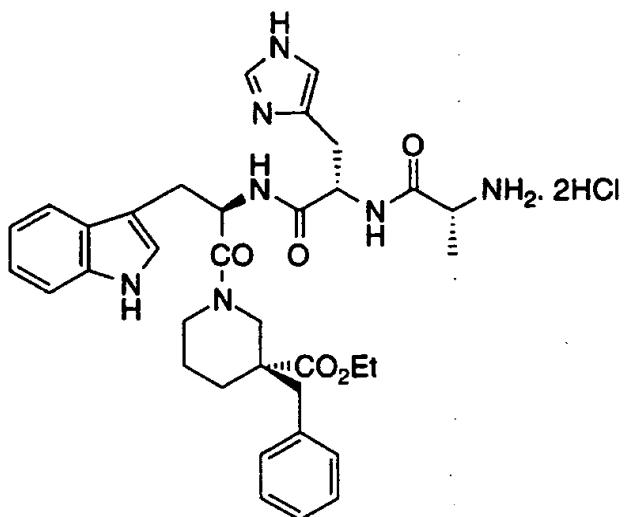
20 was stirred at room temperature overnight and then partitioned between EtOAc/H₂O. The organic phase was separated, washed with 10% NaHCO₃, water, 10% citric acid, water, then dried over MgSO₄,

- 122 -

filtered and concentrated *in vacuo*. The crude product thus obtained was purified by flash chromatography (20:1 CH₂Cl₂/MeOH) to provide 81 mg of the Boc-protected tripeptide. The protected tripeptide (81 mg) was then reacted at room temperature with a cold solution of saturated 5 HCl/THF (5 ml) for 1h. The solvent was removed *in vacuo*, and the residue was triturated with dry ether to provide the titled compound as a white solid (70 mg). FAB-MS: m/e calculated for C₃₅H₄₃N₇O₅: 641; found: 642 (M+H).

10

EXAMPLE 34



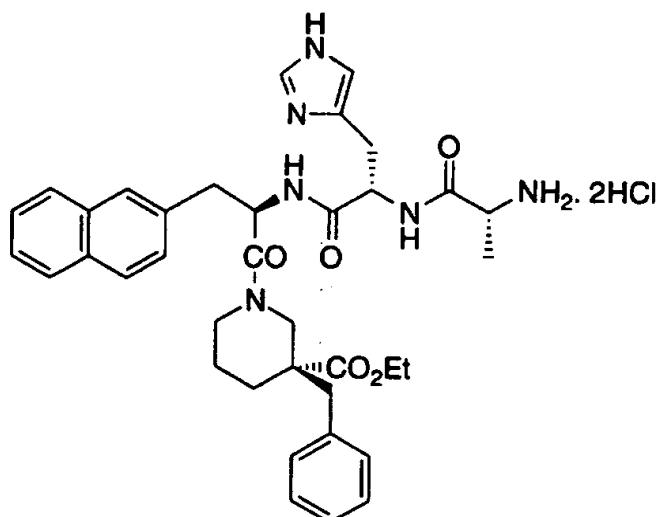
To a cold (0°C) solution of the titled compound from Example 33, Step A (126 mg) in CH₂Cl₂ (2 ml) were added Boc-D-Ala (47 mg), HOBT (50 mg), N-methylmorpholine (49 ml) and EDC (58 mg). The reaction was stirred at room temperature overnight, and then partitioned between EtOAc/H₂O layers. The organic phase was separated and washed with 10% NaHCO₃, water, 10% citric acid, water, then dried over MgSO₄, filtered and concentrated *in vacuo* to give the 15 crude product. Purification of the crude product by flash chromatography (20:1 CH₂Cl₂/MeOH) provided 85 mg of the Boc-protected tripeptide. The protected tripeptide (85 mg) was then reacted with a cold solution of saturated HCl/THF (5 ml) for 1h at room 20

- 123 -

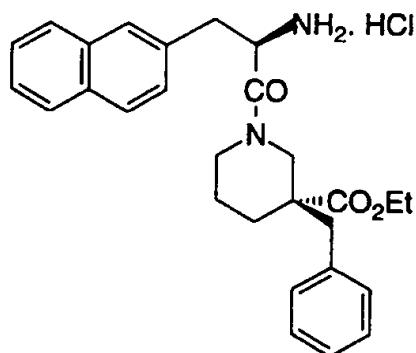
temperature. The solvent was removed in vacuo, and the residue was triturated with dry ether to provide the titled compound as a white solid (69 mg). FAB-MS: m/e calculated for C₃₅H₄₃N₇O₅: 641; found: 642 (M+H).

5

EXAMPLE 35



Step A:

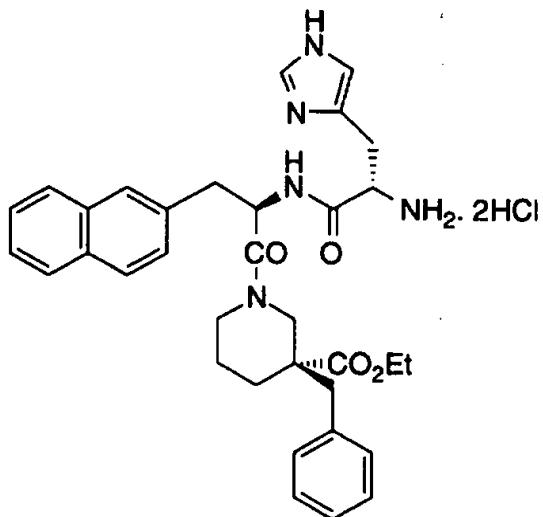


10 To a solution of Boc-D- β -Nal (0.41g) in CH₂Cl₂ (5 ml) at
 0°C were added ethyl-3-(S)-benzyl-nipecotate (0.36 g) (obtained from
 Example 28, Step C), HOBT (0.282 g), and EDC (0.296 g). The
 mixture was stirred at room temperature overnight and then partitioned
 between EtOAc/water. The organic phase was separated and washed

- 124 -

with 10% NaHCO₃, water, 10% citric acid, water, then dried over MgSO₄, filtered and concentrated *in vacuo*. The crude product, thus obtained, was purified by flash column chromatography on silica-gel using EtOAc-hexanes (1:3) as the eluent to provide the desired Boc-D-b-5 Nal amide (0.6 g). The protected intermediate was then dissolved in a saturated solution of HCl(g) in EtOAc (6 ml), and the solution was stirred at room temperature for 1 hour. The solvent and excess reagent was removed *in vacuo*, and the solid residue obtained was triturated with dry ether, filtered and dried in *vacuo* over P₂O₅ overnight to give 10 the titled product (0.5 g). FAB-MS: m/e calculated for C₂₈H₃₂N₂O₃: 444; found: 445 (M+H).

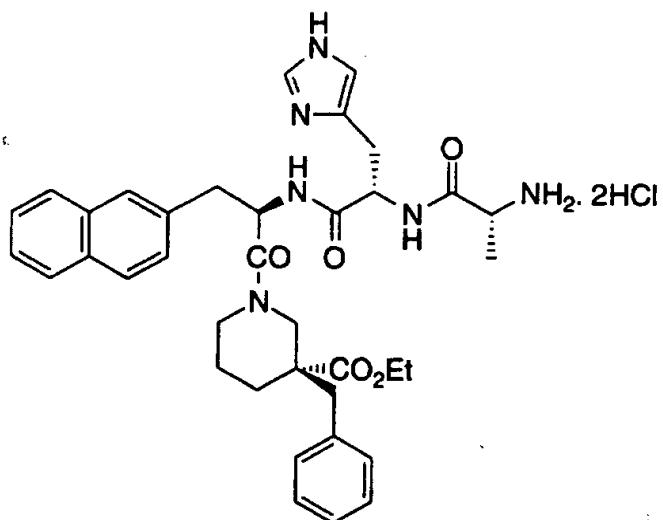
Step B:



15 The title compound was prepared from the intermediate obtained in Step A (0.2 g) using the procedure described in Example 33, Step A. FAB-MS: m/e calculated for C₃₄H₃₉N₅O₄: 581; found: 582 (M+H).

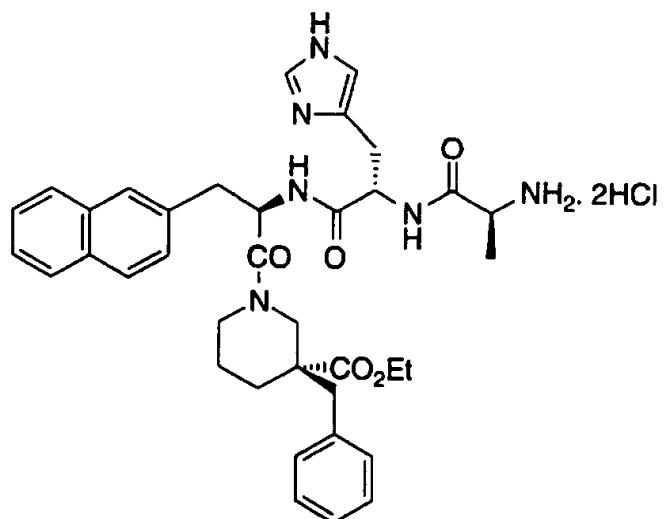
- 125 -

Step C:



The titled compound was prepared from the intermediate obtained by using the procedure described in Example 34. FAB-MS: m/e calculated for $C_{37}H_{44}N_6O_5$: 652; found: 653 ($M+H$).

EXAMPLE 36

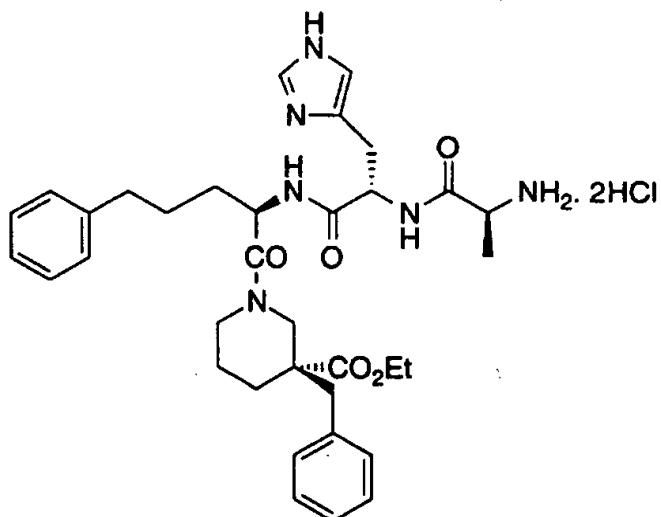


The titled compound was prepared from the intermediate obtained in Example 35, Step B using the procedure described in

- 126 -

Example PKC/JM 1, Step B. FAB-MS: m/e calculated for C₃₇H₄₄N₆O₅: 652; found: 653 (M+H).

EXAMPLE 37

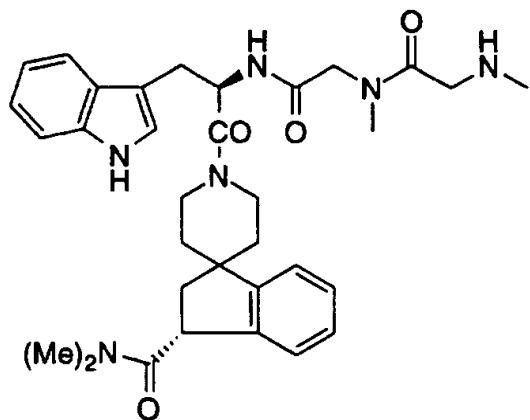


5

The titled compound was prepared by the procedure described in Example 35, using (D)-N-t-Boc-2-amino-5-phenylpentanoic acid instead of Boc-D-β-Nal in Step A. FAB-MS: m/e calculated for C₃₅H₄₆N₆O₅: 630; found: 631 (M+H).

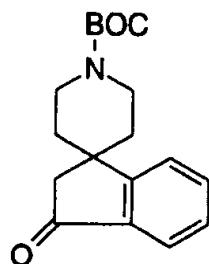
10

EXAMPLE 38



- 127 -

Step A:



To a solution of 51.0 g (0.177 mol) of 1'-(t-butyloxycarbonyl)spiro[1H-indene-1,4'-piperidine] [prepared by the method of Chambers, et al, *J. Med. Chem.*, 1992, 35, 2036] in 200 ml of THF was added 430 ml (0.5 M in THF, 0.213 mol) of 9-BBN. The reaction mixture was heated at 70°C until TLC analysis indicated that the starting material was consumed (18 hrs). The solution was concentrated to ~300 ml and then cooled to 0°C and quenched with methanol (10 ml). 4 N

5 Sodium hydroxide (213 ml) and 30% hydrogen peroxide (108 ml) were added via an addition funnel over 45 minutes. The reaction mixture was stirred for 3.5 hours and then solid sodium sulfite was added until starch paper indicated that no more peroxides were present. The reaction mixture was extracted with ethyl acetate (4 X 1 vol). The ethyl acetate

10 layer was dried over magnesium sulfate filtered and concentrated. The crude material was dissolved in dichloromethane (300 ml) and the solution was cooled to 0°C then celite (25 g) and PCC (57 g) were added in five portions over 20 minutes. The reaction mixture was warmed to room temperature and stirred overnight. The solution was then diluted

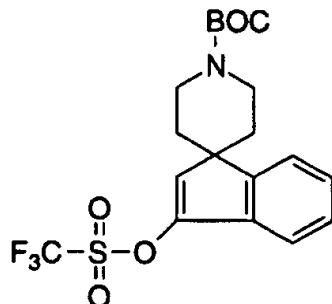
15 with ether and filtered through a pad of a mixture of celite and florisil. Purification by flash chromatography (silica gel, hexane/ethyl acetate, 5:1 to 3:1) gave 58.6 g of the title compound. ^1H NMR (200 MHz, CDCl_3): 7.75-7.60 (m, 2H), 7.50-7.44 (m, 2H), 4.30-4.15 (m, 2H), 2.85 (dt, 2H), 2.63 (s, 2H), 1.98 (dt, 2H), 1.53-1.40 (m, 2H), 1.49 (s, 9H).

20

25

- 128 -

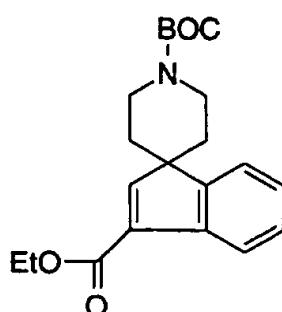
Step B:



Potassium bis(trimethylsilyl)amide (127.5 ml, 0.5 M) was added to the ketone (16.0 g, 53 mmol) in THF (200 mL) at 0°C. The 5 reaction mixture was stirred for one hour and then N-phenyltrifluoromethanesulfonamide was added. The ice bath was allowed to melt and the reaction mixture was stirred overnight at room temperature. Water was added and the aqueous layer was extracted with ethyl acetate (3 X 1 vol). The organic layer was washed with brine and then dried over 10 magnesium sulfate, filtered and then concentrated. The crude product was purified by flash chromatography (hexane/ethyl acetate 8 : 1) to give the title compound (17.8 g) as a waxy solid. ^1H NMR (200 MHz, CDCl_3): 7.65-7.14 (m, 4 H), 6.66 (s, 1 H), 4.30-4.15 (m, 2 H), 3.24-2.96 (m, 2 H), 2.06 (dt, 2 H), 1.50 (s, 9 H), 1.49-1.38 (m, 2 H).

15

Step C:

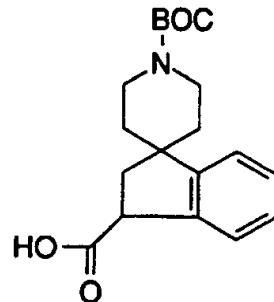


A solution of 17.4 g of the intermediate from Step B, 11.0 ml of triethylamine, 634 mg of triphenylphosphine, and 240 mg of 20 palladium acetate in 72 ml of ethanol and 158.0 ml of DMF was purged

- 129 -

for 10 minutes with carbon monoxide and then stirred under a carbon monoxide atmosphere for 24 hours. The ethanol was removed in vacuum and the reaction mixture was diluted with water and extracted repeatedly with ethyl acetate. The ethyl acetate layer was washed with 5 1N HCl, water, and brine and then dried over magnesium sulfate, filtered and concentrated. Purification by flash chromatography (hexane / ethyl acetate 8:1) provided 27.6 g of the title compound as a colorless oil. ^1H NMR (200 MHz, CDCl_3): 8.0-7.94 (m, 1H), 7.7 (s, 1 H), 7.4-7.25 (m, 3H), 4.4 (q, 2H), 4.25-4.15 (m, 2H), 3.13 (dt, 2H), 2.03 10 (dt, 2H), 1.5 (s, 9H), 1.55-1.35 (m, 2H), 1.4 (t, 3H).

Step D:



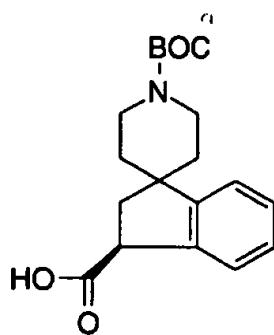
To a suspension of Pd/C (1.7g) in ethanol (300 ml) was 15 added the title compound (27 g) from Step C. The reaction mixture was purged with hydrogen and then shaken under a hydrogen atmosphere for 3 hours. The mixture was purged with nitrogen and filtered through celite and concentrated to give the title compound (27 g). The crude product was dissolved in ethanol (200 ml) and 2N sodium 20 hydroxide (76 ml) was added. The reaction mixture was heated to 50°C for three hours then cooled and the ethanol was removed under vacuum and the residue was dissolved in ethyl acetate. 1N HCl was added and the layers were separated and the aqueous layer was extracted with ethyl acetate (3 x 1 vol). The combined organic layers were washed with 25 saturated aqueous NaCl, dried over anhydrous sodium sulfate, filtered and concentrated to provide the title compound (23.8 g) as a white solid. ^1H NMR (200 MHz, CDCl_3): 7.50-7.42 (m, 1 H), 7.34-7.12 (m, 3 H),

- 130 -

4.22-4.04 (m, 3 H), 3.06-2.84 (m, 2 H), 2.40 (d, 2 H), 1.88-1.6 (m, 4 H), 1.50 (s, 9 H).

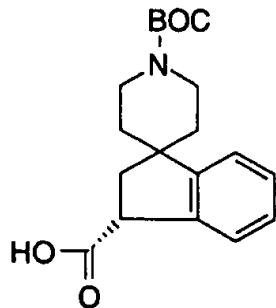
Step E:

5



The acid from Step D (23.5 g, 0.07 mol) was dissolved in toluene (150 ml) and R-methylbenzylamine (9.02 ml) was added. The toluene solution was heated on a steam bath until everything was in solution. The solution was then seeded with crystals grown in the same 10 way on a much smaller scale. The solution was allowed to sit overnight and then the mixture was filtered to give 15.8 g of crystals. The crystals were recrystallized from toluene two more times. The crystals (12 g) were dissolved in ethyl acetate /1 N HCl and the organic layer was washed with 1 N HCl (2 X 1 vol) and brine. The organic layer 15 was dried over magnesium sulfate, filtered and concentrated to give 8.9 g of the title compound. $[\alpha]^D = -16.9$ (c= 0.84, methanol) .

Step F:

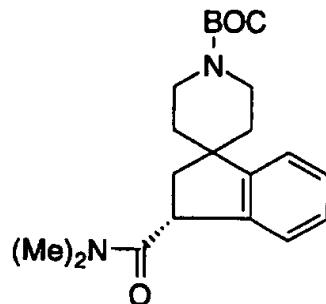


- 131 -

The mother liqueurs from Step E were washed with 1 N HCl (2 x 1 vol) and brine dried over magnesium sulfate filtered and concentrated to give recovered acid (15.4 g). To this acid in toluene (100 mL) was added S-methylbenzylamine (5.95 mL). The crystals 5 were recrystallized four times from toluene as above to give 12.3 g of salt. The salt was dissolved in ethyl acetate / 1 N HCl and washed with 1 N HCl (2 X 1 vol) and brine. The organic layer was dried over magnesium sulfate and filtered and concentrated to give the title compound (9.0 g). $[\alpha]_D = 17.1$ (c=1.06, methanol).

10

Step G:



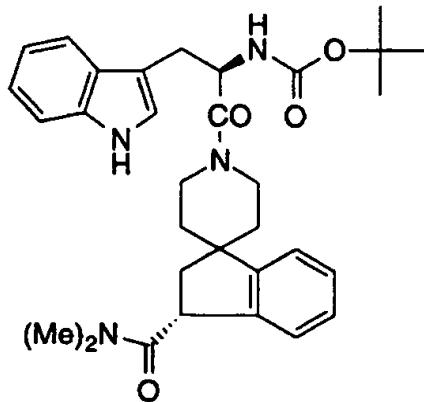
A solution of the title compound from Step F (14.0 g, 42 mMol) in dichloromethane was cooled to 0° and dimethylamine (25.4 mL, 2M in THF) was added. The mixture was stirred for ten minutes 15 at 0°C and then EDC and DMAP were added. The reaction mixture was stirred for four hours at 0°C and then quenched with 1 N HCl. The aqueous layers were extracted with dichloromethane and the combined organic layers were then washed with water and brine and dried over sodium sulfate. The crude product was purified by flash 20 chromatography (dichloromethane/acetone 9:1) to give the title compound (12.2 g). HPLC analysis (chiralcel OD-R, 50% 0.5N NaClO₄ / 50% acetonitrile, 0.5 ml/min. E₁ retention time 20.8 min (E₁ prepared from the intermediate in Example 38 Step E as in Example 38 25 Step G; E₂ retention time 24.7 min) showed it to be approximately a 1 : 200 mixture of enantiomers. ¹HNMR (400 MHz, CDCl₃): 7.25-7.05(m, 4 H), 4.35 (t, 1 H), 4.20-4.10 (m, 2 H), 3.25 (s, 3 H), 3.05 (s, 3 H), 2.90-

- 132 -

2.85 (m, 2H), 2.42-2.28 (m, 2H), 1.95 (dt, 1H), 1.75-1.60 (m, 2 H), 1.52-1.50 (m, 1 H), 1.49 (s, 9H).

Step H:

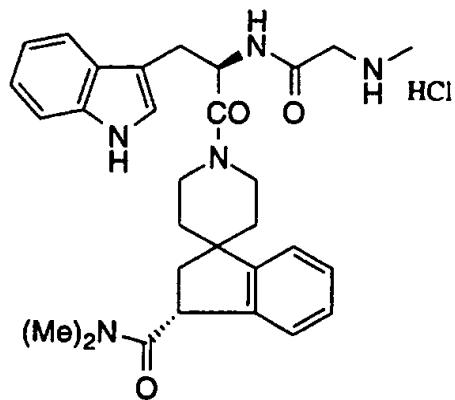
5



The title compound from Step G (6.4 g, 18.4 mmol) was stirred in ethyl acetate saturated with HCl for two hours and then concentrated and azeotroped from dichloromethane (2X) and toluene (1X). The residue was dissolved in dichloromethane, cooled to 0°C and 10 Boc-D-tryptophan (6.2 g, 20.2 mMol), NMM (2.0 mL, 18.4 mMol), HOBT (3.7 g, 27.6 mMol) and finally EDC (5.27 g, 27.6 mMol) were added. The reaction mixture was stirred at room temperature overnight and then poured into ethyl acetate. The organic layer was washed with saturated bicarb, 1 N HCl, water and finally brine. The organic layers 15 were dried over magnesium sulfate, filtered and concentrated. Purification by flash chromatography (ethyl acetate) gave the title compound (4.6 g).

- 133 -

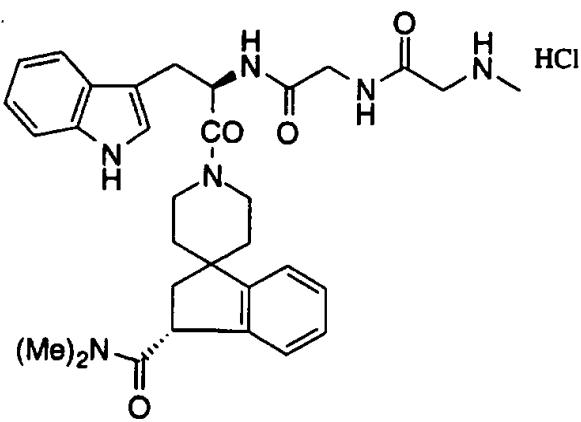
Step I:



The title compound from Step H (111 mg, 0.230 mMol) was deprotected with ethyl acetate / HCl as above. The residue was 5 coupled to Boc-sarcosine using the procedure for Step H. The reaction mixture was loaded directly onto a flash column and eluted with dichloromethane/ acetone (1 : 1) to give the Boc protected product (122 mg). This material was deprotected as above with ethyl acetate / HCl to give the title compound (120 mg).

10

Step J:



The title compound from Step J (40 mg 0.074 mMol) was coupled to Boc-sarcosine as in Step I. Purification of the crude product 15 by flash chromatography (dichloromethane / acetone 1 : 1) gave the Boc

- 134 -

protected title compound (43 mg). Removal of the Boc group with ethyl acetate / HCl followed by purification by MPLC (LH₂O, methanol) gave the title compound (34 mg). ESI-MS: 587.3 (M+1).

5

While the invention has been described and illustrated with reference to certain particular embodiments thereof, those skilled in the art will appreciate that various adaptations, changes, modifications, substitutions, deletions, or additions of procedures and protocols may be made without departing from the spirit and scope of the invention. For example, effective dosages other than the particular dosages as set forth herein above may be applicable as a consequence of variations in the responsiveness of the mammal being treated for any of the indications with the compounds of the invention indicated above. Likewise, the specific pharmacological responses observed may vary according to and depending upon the particular active compounds selected or whether there are present pharmaceutical carriers, as well as the type of formulation and mode of administration employed, and such expected variations or differences in the results are contemplated in accordance with the objects and practices of the present invention. It is intended, therefore, that the invention be defined by the scope of the claims which follow and that such claims be interpreted as broadly as is reasonable.

10

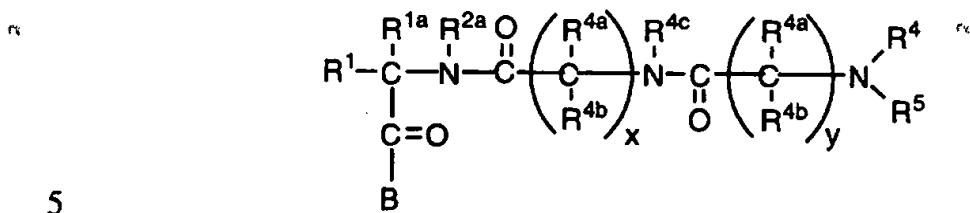
15

20

- 135 -

WHAT IS CLAIMED IS:

1. A compound of the formula:



wherein:

R¹ is selected from the group consisting of:
 C₁-C₁₀ alkyl-, -aryl-, aryl (C₁-C₆ alkyl)-,
 10 heteroaryl-, heteroaryl(C₁-C₆ alkyl)-,
 (C₃-C₇ cycloalkyl)-(C₁-C₆ alkyl)-,
 (C₁-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 aryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 heteroaryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-, and
 15 (C₃-C₇ cycloalkyl)-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 wherein K is -O-, -S(O)_m-, -N(R²)C(O)-, -C(O)N(R²)-, -OC(O)-,
 -C(O)O-, -CR²=CR²- or -C≡C-,
 wherein R² and the alkyl groups may be further substituted with 1 to 9
 halo, -S(O)_mR^{2a}, 1 to 3 of -OR^{2a}, or -C(O)OR^{2a}, and wherein aryl is
 20 phenyl or naphthyl, and heteroaryl is selected from indolyl, thiophenyl,
 benzofuranyl, benzothiophenyl, aza-indolyl, pyridinyl, quinolinyl, and
 benzimidazolyl, wherein aryl and heteroaryl are unsubstituted or
 substituted with phenyl, phenoxy, halophenyl, 1 to 3 of -C₁-C₆ alkyl, 1
 to 3 of halo, 1 to 2 of -OR², methylenedioxy, -S(O)_mR², 1 to 2 of
 25 -CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²), -C(O)OR²,
 -C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, or -N(R²)SO₂R²;

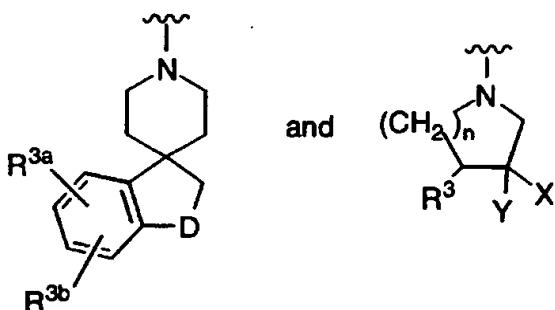
R^{1a} is hydrogen or C₁-C₄ alkyl;

- 136 -

R^2 is selected from the group consisting of:
 hydrogen, $-C_1-C_6$ alkyl, $-C_3-C_7$ cycloalkyl, and $-CH_2$ -phenyl,
 wherein the alkyl or the cycloalkyl is unsubstituted or substituted with
 hydroxyl, C_1-C_3 alkoxy, thioalkyl, $C(O)OR^{2a}$, and where, if two $-C_1$ -
 5 C_6 alkyl groups are present on one atom, they may be joined to form a
 C_3-C_8 cyclic ring being selected from the group consisting of
 pyrrolidine, piperidine, piperazine, morpholine, thiomorpholine,
 optionally substituted by hydroxyl;

10 R^{2a} is hydrogen or C_1-C_6 alkyl;

B is selected from:



R^3 is selected from: hydrogen, $-(CH_2)_r$ phenyl, $-(CH_2)_r$ thienyl,
 15 $-(CH_2)_r$ benzimidazolyl, $-(CH_2)_r$ quinolinyl, $-(CH_2)_r$ naphthyl,
 $-(CH_2)_r$ indolyl, $-C_1-C_{10}$ alkyl, $-C_3-C_7$ cycloalkyl, where the phenyl,
 naphthyl, indolyl, thienyl, benzimidazolyl, quinolinyl, and C_3-C_7
 cycloalkyl rings may be substituted by 1 to 3 substituents selected from
 the group consisting of: C_1-C_6 alkyl, halogen, $-OR^2$, $-NHSO_2CF_3$,
 20 $-(CH_2)_rOR^6$, $-(CH_2)_rN(R^2)(R^6)$, $-(CH_2)_r(R^6)$, $-(CH_2)_rC(O)OR^2$,
 $-(CH_2)_rC(O)OR^6$, $-(CH_2)_rOC(O)R^2$, $-(CH_2)_rOC(O)R^6$,
 $-(CH_2)_rC(O)R^2$, $-(CH_2)_rC(O)R^6$, $-(CH_2)_rC(O)N(R^2)(R^2)$,
 $-(CH_2)_rC(O)N(R^2)(R^6)$, $-(CH_2)_rN(R^2)C(O)(R^2)$,
 $-(CH_2)_rN(R^2)C(O)R^6$, $-(CH_2)_rN(R^6)C(O)R^2$, $-(CH_2)_rN(R^6)C(O)R^6$,
 25 $-(CH_2)_rN(R^2)C(O)OR^2$, $-(CH_2)_rN(R^2)C(O)OR^6$,
 $-(CH_2)_rN(R^6)C(O)OR^2$, $-(CH_2)_rN(R^6)C(O)OR^6$,
 $-(CH_2)_rN(R^2)C(O)N(R^2)(R^6)$, $-(CH_2)_rN(R^2)C(O)N(R^2)(R^2)$,

- 137 -

-(CH₂)_rN(R⁶)C(O)N(R²)(R⁶), -(CH₂)_rN(R²)SO₂R²,
-(CH₂)_rN(R⁶)SO₂R², -(CH₂)_rN(R⁶)SO₂R⁶, -(CH₂)_rOC(O)N(R²)(R⁶),
-(CH₂)_rOC(O)N(R²)(R²), -(CH₂)_rSO₂N(R²)(R⁶),
-(CH₂)_rOC(O)N(R²)(R²), -(CH₂)_rSO₂N(R²)(R⁶),
5 -(CH₂)_rSO₂N(R²)(R²), -(CH₂)_rN(R²)SO₂N(R²)(R⁶),
-(CH₂)_rN(R⁶)SO₂N(R²)(R⁶), -(CH₂)_rS(O)_mR⁶, and -(CH₂)_rS(O)_mR²;

R^{3a} and R^{3b} are independently selected from: hydrogen, phenyl,
phenoxy, halophenyl, -C₁-C₆ alkyl, halogen, -OR², methylenedioxy,
10 -S(O)_mR², -CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²),
-C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, and
-N(R²)SO₂R²;

R⁴ and R⁵ are independently selected from hydrogen, C₁-C₆ alkyl, and
15 substituted C₁-C₆ alkyl where the substituents are selected from halo,
hydroxy, phenyl, and C₁-C₆ alkoxy carbonyl;
or R⁵ and R⁴ may be taken together to form -(CH₂)_d-La(CH₂)_e- where
La is -C(R²)₂-, -O-, -S(O)_m- or -N(R²)-, d and e are independently 1 to
3 and R² is as defined above;

20 R^{4a} and R^{4b} are independently selected from: hydrogen, C₁-C₆ alkyl,
trifluoromethyl, phenyl, or substituted C₁-C₆ alkyl where the
substituents are selected from: imidazolyl, naphthyl, phenyl, indolyl,
p-hydroxyphenyl, -OR², -S(O)_mR², -C(O)OR², C₃-C₇ cycloalkyl,
25 -N(R²)(R²), -C(O)N(R²)(R²); or R^{4a} and R^{4b} may independently be
joined to one or both of R⁴ or R^{4c} to form an alkylene bridge between
the terminal nitrogen and the alkyl portion of the R^{4a} or R^{4b} and the
R⁴ or R^{4c} group, wherein the bridges contain 1 to 8 carbons atoms; or
R^{4a} and R^{4b} may be joined to one another to form C₃-C₇ cycloalkyl;

30 R^{4c} is selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆
alkyl where the substituents are selected from 1 to 5 halo, 1 to 3
hydroxy, phenyl, naphthyl, imidazolyl, indolyl, and C₁-C₆
alkoxy carbonyl;

R^6 is selected from: hydrogen, C₁-C₆ alkyl, and (CH₂)_qaryl, wherein the (CH₂)_q and alkyl groups may be optionally substituted by -O(R²), -S(O)_mR², -C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), or

5 -N(R²)C(O)N(R²)(R²), wherein the aryl group is selected from: phenyl, pyridyl, 1H-tetrazolyl, triazolyl, oxadiazolyl, pyrazolyl, thiadiazolyl, and benzimidazol-2-yl, which is optionally substituted with C₁-C₆ alkyl, C₃-C₆ cycloalkyl, amino, or hydroxyl;

10 X is selected from the group consisting of: hydrogen, -C≡N, -(CH₂)_qN(R²)C(O)R², -(CH₂)_qN(R²)C(O)(CH₂)_taryl, -(CH₂)_qN(R²)SO₂(CH₂)_taryl, -(CH₂)_qN(R²)SO₂R², -(CH₂)_qN(R²)C(O)N(R²)(CH₂)_taryl, -(CH₂)_qN(R²)C(O)N(R²)(R²), -(CH₂)_qC(O)N(R²)(R²), -(CH₂)_qC(O)N(R²)(CH₂)_taryl,

15 -(CH₂)_qC(O)OR², -(CH₂)_qC(O)O(CH₂)_taryl, -(CH₂)_qOR², -(CH₂)_qOC(O)R², -(CH₂)_qOC(O)(CH₂)_taryl, -(CH₂)_qOC(O)N(R²)(R²), -(CH₂)_qC(O)R², -(CH₂)_qC(O)(CH₂)_taryl, -(CH₂)_qN(R²)C(O)OR², -(CH₂)_qN(R²)SO₂N(R²)(R²), -(CH₂)_qS(O)_mR², and -(CH₂)_qS(O)_m(CH₂)_taryl, where R², (CH₂)_q

20 and (CH₂)_t group may be optionally substituted with C₁-C₄ alkyl, hydroxyl, C₁-C₄ lower alkoxy, carboxyl, N(R²)(R²), CONH₂, S(O)_mCH₃, carboxylate C₁-C₄ alkyl esters, or 1H-tetrazol-5-yl, and aryl is phenyl, naphthyl, pyridyl, thiazolyl, or 1H-tetrazol-5-yl groups which may be optionally substituted with halogen, -OR², -CON(R²)(R²), -C(O)OR², C₁-C₄ alkyl, -S(O)_mR², or 1H-tetrazol-5-yl;

25 Y is selected from the group consisting of: hydrogen, C₁-C₁₀ alkyl, -(CH₂)_taryl, -(CH₂)_q(C₃-C₇ cycloalkyl), -(CH₂)_q-K-(C₁-C₆ alkyl), -(CH₂)_q-K-(CH₂)_taryl, -(CH₂)_q-K-(CH₂)_t(C₃-C₇ cycloalkyl containing O, NR² S) and -(CH₂)_q-K-(CH₂)_t(C₃-C₇ cycloalkyl), where K is O, S(O)_m, C(O)NR², CH=CH, C≡C, N(R²)C(O), C(O)NR², C(O)O, or OC(O), and where the alkyl, R², (CH₂)_q and (CH₂)_t groups are optionally

- 139 -

substituted by C₁-C₄ alkyl, hydroxyl, C₁-C₄ lower alkoxy, carboxyl, -CONH₂ or a carboxylate C₁-C₄ alkyl ester, and aryl is phenyl, naphthyl, pyridyl, 1-H-tetrazol-5-yl, thiazolyl, imidazolyl, indolyl, oxadiazolyl, pyrimidinyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl, 5 thiophenyl, quinolinyl, pyrazinyl, or isothiazolyl which is optionally substituted with halogen, -OR², -C(O)OR², N(R²)(R²), -C(O)N(R²)(R²), nitro, cyano, benzyl, C₁-C₄ alkyl, -S(O)_mR², or 1H-tetrazol-5-yl;

10 D is selected from: -N(R⁷)-, -S(O)_m-, -C(O)- and -C(H)(R⁷)-, wherein R⁷ is selected from: -R², -OR², -(CH₂)_qaryl, -C(O)R², -C(O)(CH₂)_qaryl, -SO₂R², -SO₂(CH₂)_qaryl, -C(O)N(R²)(R²), -C(O)N(R²)(CH₂)_qaryl, -C(O)OR², 1-H-tetrazol-5-yl, -SO₂N(R²)aryl, -SO₂N(R²)(R²) and the (CH₂)_q may be optionally substituted by C₁-C₄ 15 alkyl, and the R² and aryl may be optionally further substituted with a substituent selected from: -OR^{2a}, -O(CH₂)_qaryl, -C(O)OR^{2a}, -C(O)(CH₂)_qaryl, -C(O)N(R^{2a})(R^{2a}), -C(O)N(R^{2a})(CH₂)₁aryl, halogen, -N(R^{2a})(R^{2a}), -C₁-C₄ alkyl, 1,2,4-triazolyl, 1-H-tetrazol-5-yl, -C(O)NHSO₂R^{2a}, -S(O)_mR^{2a}, -C(O)NHSO₂(CH₂)_qaryl, 20 -N(R²)C(O)N(R^{2a})(R^{2a}), -N(R^{2a})C(O)N(R^{2a})(CH₂)_qaryl, -N(R^{2a})(R^{2a}), -N(R^{2a})C(O)R^{2a}, -N(R^{2a})C(O)(CH₂)_qaryl, -OC(O)N(R^{2a})(R^{2a}), -OC(O)N(R^{2a})(CH₂)_qaryl;

25 1 is 0, 1 or 2;

25 m is 0, 1, or 2;

n is 0, 1, or 2;

q is 0, 1, 2, 3, or 4;

r is 0, 1, 2, or 3;

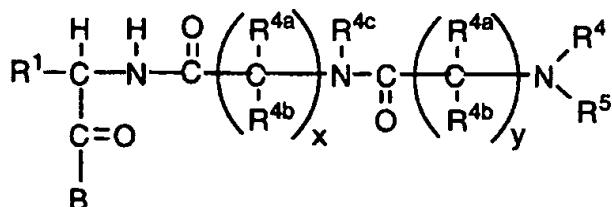
t is 0, 1, 2, or 3;

30 v is 0, 1, or 2;

and pharmaceutically acceptable salts and individual diastereomers thereof.

- 140 -

2. The compound of Claim 1 of the formula:



5 wherein:

R¹ is selected from the group consisting of:
 C₁-C₁₀ alkyl, -aryl-, aryl (C₁-C₆ alkyl)-,
 heteroaryl-, heteroaryl(C₁-C₆ alkyl)-,
 (C₃-C₇ cycloalkyl)-(C₁-C₆ alkyl)-,

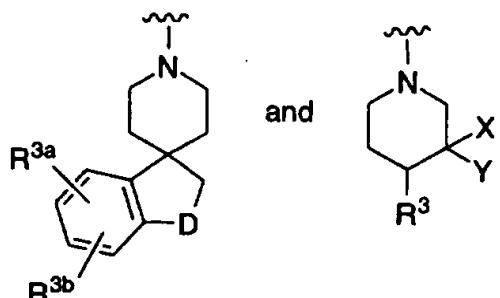
10 (C₁-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 aryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 heteroaryl-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-, and
 (C₃-C₇ cycloalkyl)-(C₀-C₅ alkyl)-K-(C₁-C₅ alkyl)-,
 wherein K is -O-, -S(O)_m-, -N(R²)C(O)-, -C(O)N(R²)-, -OC(O)-,
 15 -C(O)O-, -CR²=CR²- or -C≡C-,
 wherein R² and the alkyl groups may be further substituted with 1 to 9
 halo, -S(O)_mR^{2a}, 1 to 3 of -OR^{2a}, or -C(O)OR^{2a}, and wherein aryl is
 phenyl or naphthyl, and heteroaryl is selected from indolyl, thiophenyl,
 benzofuranyl, benzothiophenyl, aza-indolyl, pyridinyl, quinolinyl, and
 20 benzimidazolyl, wherein aryl and heteroaryl are unsubstituted or
 substituted with phenyl, phenoxy, halophenyl, 1 to 3 of -C₁-C₆ alkyl, 1
 to 3 of halo, 1 to 2 of -OR², methylenedioxy, -S(O)_mR², 1 to 2 of
 -CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²), -C(O)OR²,
 -C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, or -N(R²)SO₂R²;

25 R² is selected from the group consisting of:
 hydrogen, -C₁-C₆ alkyl, -C₃-C₇ cycloalkyl, and -CH₂-phenyl,
 wherein the alkyl or the cycloalkyl is unsubstituted or substituted with
 hydroxyl, C₁-C₃ alkoxy, thioalkyl, -C(O)OR^{2a}, and wherein, if two
 30 -C₁-C₆ alkyl groups are present on one atom, the groups may be
 optionally joined to form a C₃-C₈ cyclic ring being selected from the

- 141 -

group consisting of pyrrolidine, piperidine, piperazine, morpholine, thiomorpholine;

B is selected from:



5

R^3 is selected from: hydrogen, phenyl, naphthyl, indolyl, benzimidazolyl, thienyl, quinolinyl, where the phenyl, naphthyl, benzimidazolyl, thienyl, quinolinyl, and indolyl may be substituted by 1 to 3 substituents selected from the group consisting of: C₁-C₆ alkyl, halogen, -OR², -(CH₂)_rOR⁶, -(CH₂)_rN(R²)(R⁶), -(CH₂)_r(R⁶), -(CH₂)_rC(O)OR², -(CH₂)_rC(O)OR⁶, -(CH₂)_rC(O)R², -(CH₂)_rC(O)R⁶, -(CH₂)_rC(O)N(R²)(R²), -(CH₂)_rC(O)N(R²)(R⁶), -(CH₂)_rN(R²)C(O)(R²), -(CH₂)_rN(R²)C(O)R⁶, -(CH₂)_rN(R⁶)C(O)R², -(CH₂)_rN(R⁶)C(O)R⁶, -(CH₂)_rN(R²)C(O)OR², 10 -(CH₂)_rN(R²)C(O)OR⁶, -(CH₂)_rN(R⁶)C(O)OR², -(CH₂)_rN(R⁶)C(O)OR⁶, -(CH₂)_rN(R²)C(O)N(R²)(R⁶), -(CH₂)_rN(R²)C(O)N(R⁶)(R⁶), -(CH₂)_rN(R²)SO₂R², -(CH₂)_rN(R⁶)SO₂R², -(CH₂)_rN(R⁶)SO₂R⁶, 15 -(CH₂)_rOC(O)N(R²)(R⁶), -(CH₂)_rSO₂N(R²)(R⁶), -(CH₂)_rSO₂N(R²)(R²), -(CH₂)_rS(O)_mR⁶, and 20 -(CH₂)_rS(O)_mR²;

R^{3a} and R^{3b} are independently selected from: hydrogen, phenyl, phenoxy, halophenyl, -C₁-C₆ alkyl, halogen, -OR², methylenedioxy, 25 -S(O)_mR², -CF₃, -OCF₃, nitro, -N(R²)(R²), -N(R²)C(O)(R²), -C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), -N(R²)SO₂-aryl, and -N(R²)SO₂R²;

- 142 -

R⁴ and R⁵ are independently selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from halo, hydroxy, phenyl, and C₁-C₆ alkoxy carbonyl;

5 or R⁵ and R⁴ may be taken together to form -(CH₂)_d-L_a(CH₂)_e- where L_a is -C(R²)₂-, -O-, -S(O)_m- or -N(R²)-, d and e are independently 1 to 3 and R² is as defined above;

R^{4a} and R^{4b} are independently selected from: hydrogen, C₁-C₆ alkyl, 10 trifluoromethyl, phenyl, or substituted C₁-C₆ alkyl where the substituents are selected from: imidazolyl, naphthyl, phenyl, indolyl, p-hydroxyphenyl, -OR², -S(O)_mR², -C(O)OR², C₃-C₇ cycloalkyl, -N(R²)(R²), -C(O)N(R²)(R²); or R^{4a} and R^{4b} may independently be joined to one or both of R⁴ or R^{4c} to form an alkylene bridge between 15 the terminal nitrogen and the alkyl portion of the R^{4a} or R^{4b} and the R⁴ or R^{4c} group, wherein the bridges contain 1 to 5 carbons atoms; or R^{4a} and R^{4b} may be joined to one another to form C₃-C₇ cycloalkyl;

R^{4c} is selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ 20 alkyl where the substituents are selected from 1 to 5 halo, 1 to 3 hydroxy, phenyl, C₁-C₆ alkoxy carbonyl;

R⁶ is selected from: hydrogen, C₁-C₆ alkyl, and (CH₂)_varyl, wherein 25 the (CH₂)_v and alkyl groups may be optionally substituted by -O(R²), -S(O)_mR², -C(O)OR², -C(O)N(R²)(R²), -SO₂N(R²)(R²), or -N(R²)C(O)N(R²)(R²), wherein the aryl group is selected from: phenyl, pyridyl, 1H-tetrazolyl, triazolyl, oxadiazolyl, pyrazolyl, thiadiazolyl, and benzimidazol-2-yl, which is optionally substituted with C₁-C₆ alkyl, C₃-C₆ cycloalkyl, amino, or hydroxyl;

- 143 -

X is selected from the group consisting of: hydrogen, $-C\equiv N$, $-(CH_2)_q N(R^2)C(O)R^2$, $-(CH_2)_q N(R^2)C(O)(CH_2)_t aryl$, $-(CH_2)_q N(R^2)SO_2(CH_2)_t aryl$, $-(CH_2)_q N(R^2)SO_2R^2$, $-(CH_2)_q N(R^2)C(O)N(R^2)(CH_2)_t aryl$, $-(CH_2)_q N(R^2)C(O)N(R^2)(R^2)$,

5. $-(CH_2)_q C(O)N(R^2)(R^2)$, $-(CH_2)_q C(O)N(R^2)(CH_2)_t aryl$, $-(CH_2)_q C(O)OR^2$, $-(CH_2)_q C(O)O(CH_2)_t aryl$, $-(CH_2)_q OR^2$, $-(CH_2)_q OC(O)R^2$, $-(CH_2)_q OC(O)(CH_2)_t aryl$, $-(CH_2)_q OC(O)N(R^2)(R^2)$, $-(CH_2)_q C(O)R^2$, $-(CH_2)_q C(O)(CH_2)_t aryl$, $-(CH_2)_q N(R^2)C(O)OR^2$, $-(CH_2)_q N(R^2)SO_2N(R^2)(R^2)$,

10 $-(CH_2)_q S(O)mR^2$, and $-(CH_2)_q S(O)m(CH_2)_t aryl$, where R^2 , $(CH_2)_q$ and $(CH_2)_t$ group may be optionally substituted with C₁-C₄ alkyl, hydroxyl, C₁-C₄ lower alkoxy, carboxyl, $N(R^2)(R^2)$, $CONH_2$, $S(O)_mCH_3$, carboxylate C₁-C₄ alkyl esters, or 1H-tetrazol-5-yl, and aryl is phenyl, naphthyl, pyridyl, thiazolyl, or 1H-tetrazol-5-yl groups

15 which may be optionally substituted with halogen, $-OR^2$, $-CON(R^2)(R^2)$, $-C(O)OR^2$, C₁-C₄ alkyl, $-S(O)_mR^2$, or 1H-tetrazol-5-yl;

Y is selected from the group consisting of:

20 hydrogen, C₁-C₁₀ alkyl, $-(CH_2)_t aryl$, $-(CH_2)_q(C_3-C_7$ cycloalkyl), $-(CH_2)_q-K-(C_1-C_6$ alkyl), $-(CH_2)_q-K-(CH_2)_t aryl$, $-(CH_2)_q-K-(CH_2)_t(C_3-C_7$ cycloalkyl containing O, NR^2 S) and $-(CH_2)_q-K-(CH_2)_t(C_3-C_7$ cycloalkyl), where K is O, $S(O)_m$, $C(O)NR^2$, $CH=CH$, $C\equiv C$, $N(R^2)C(O)$, $C(O)NR^2$, $C(O)O$, or $OC(O)$,

25 and where the alkyl, R^2 , $(CH_2)_q$ and $(CH_2)_t$ groups are optionally substituted by C₁-C₄ alkyl, hydroxyl, C₁-C₄ lower alkoxy, carboxyl, $-CONH_2$ or a carboxylate C₁-C₄ alkyl ester, and aryl is phenyl, naphthyl, pyridyl, 1-H-tetrazol-5-yl, thiazolyl, imidazolyl, indolyl, oxadiazolyl, pyrimidinyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl,

30 thiophenyl, quinolinyl, pyrazinyl, or isothiazolyl which is optionally substituted with halogen, $-OR^2$, $-C(O)OR^2$, $N(R^2)(R^2)$, $-C(O)N(R^2)(R^2)$, nitro, cyano, benzyl, C₁-C₄ alkyl, $-S(O)_mR^2$, or 1H-tetrazol-5-yl;

- 144 -

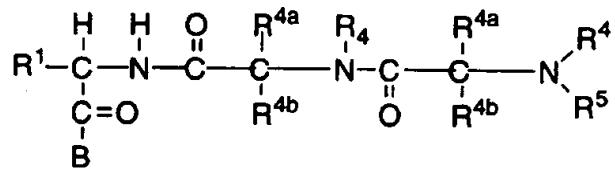
D is selected from: -N(R⁷)-, -S(O)_m-, -C(O)- and -C(H)(R⁷)-,
wherein R⁷ is selected from: -R², -(CH₂)_qaryl, -C(O)R², -SO₂R²,
-C(O)N(R²)(R²), -C(O)OR², 1-H-tetrazol-5-yl, -SO₂N(R²)aryl,
-SO₂N(R²)(R²) and the (CH₂)_q may be optionally substituted by C₁-C₄
5 alkyl, and the R² and aryl may be optionally further substituted with a
substituent selected from: -OR^{2a}, -C(O)OR^{2a}, -C(O)N(R^{2a})(R^{2a}),
halogen, -C₁-C₄ alkyl, and the aryl is selected from of triazolyl,
oxadiazolyl, thiadiazolyl, thiazolyl, imidazolyl, and 1H-tetrazolyl;

10 1 is 0, 1 or 2;
m is 0, 1, or 2;
q is 0, 1, 2, 3, or 4;
r is 0, 1, 2, or 3;
t is 0, 1, 2, or 3;
15 v is 0, 1, or 2;

and pharmaceutically acceptable salts and individual diastereomers
thereof.

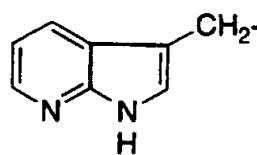
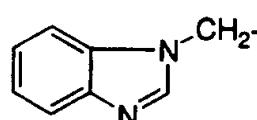
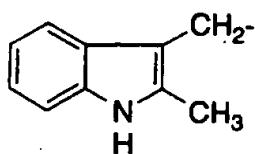
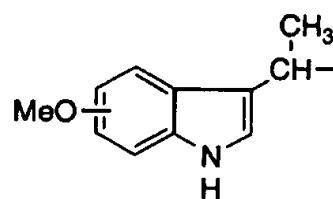
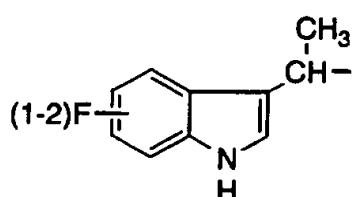
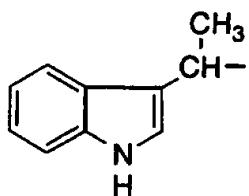
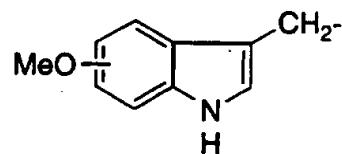
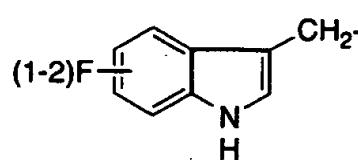
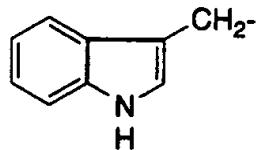
- 145 -

3. The compound of Claim 1 of the formula:

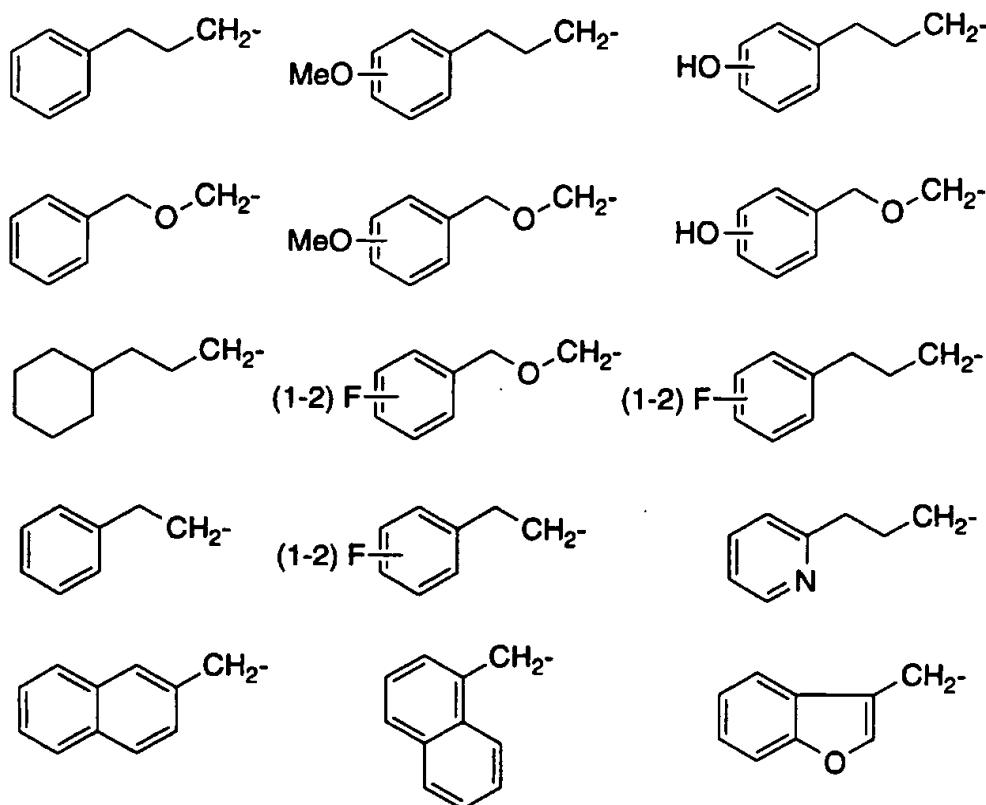


5 wherein:

R^1 is selected from the group consisting of:



- 146 -



or their regioisomers where not specified;

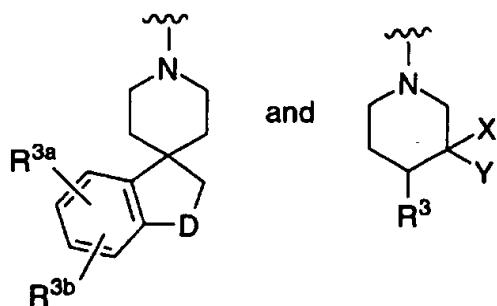
R^2 is selected from the group consisting of:

5 hydrogen, -C₁-C₆ alkyl, -C₃-C₇ cycloalkyl, and -CH₂-phenyl,
 wherein the alkyl or the cycloalkyl is unsubstituted or substituted with
 hydroxyl, C₁-C₃ alkoxy, thioalkyl, -C(O)OR^{2a}, and wherein, if two
 -C₁-C₆ alkyl groups are present on one atom, the groups may be
 optionally joined to form a C₃-C₈ cyclic ring being selected from the
 10 group consisting of pyrrolidine, piperidine, piperazine, morpholine,
 thiomorpholine;

R^{2a} is hydrogen, or C₁-C₄ alkyl;

- 147 -

B is selected from:



R³ is selected from: hydrogen or phenyl, wherein the phenyl is substituted in the ortho position by a substituent selected from the group consisting of: C₁-C₆ alkyl, halogen, -OR², -(CH₂)_rOR⁶, -(CH₂)_rN(R²)(R⁶), -(CH₂)_r(R⁶), -(CH₂)_rC(O)OR², -(CH₂)_rC(O)OR⁶, -(CH₂)_rC(O)R², -(CH₂)_rC(O)R⁶, -(CH₂)_rC(O)N(R²)(R²), -(CH₂)_rC(O)N(R²)(R⁶), -(CH₂)_rSO₂N(R²)(R⁶), -(CH₂)_rSO₂N(R²)(R²), -(CH₂)_rS(O)_mR⁶, and -(CH₂)_rS(O)_mR²;

R^{3a} and R^{3b} are independently selected from: hydrogen, -C₁-C₆ alkyl and halogen;

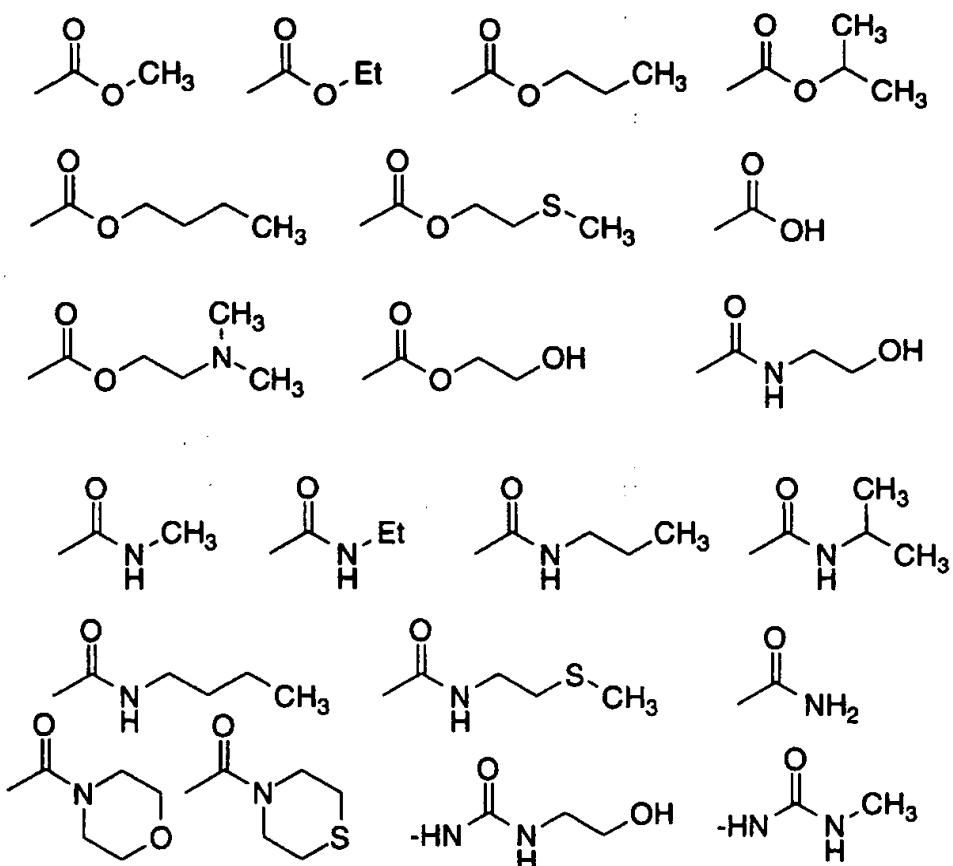
R⁴ and R⁵ are independently selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from halo, hydroxy, phenyl, and C₁-C₆ alkoxy carbonyl; or R⁵ and R⁴ may be taken together to form -(CH₂)_d-La(CH₂)_e- where La is -C(R²)₂- , -O-, -S(O)_m- or -N(R²)-, d and e are independently 1 to 3 and R² is as defined above;

R^{4a} and R^{4b} are independently selected from: hydrogen, C₁-C₆ alkyl, or substituted C₁-C₆ alkyl where the substituents are selected from: imidazolyl, naphthyl, phenyl, indolyl, and p-hydroxyphenyl;

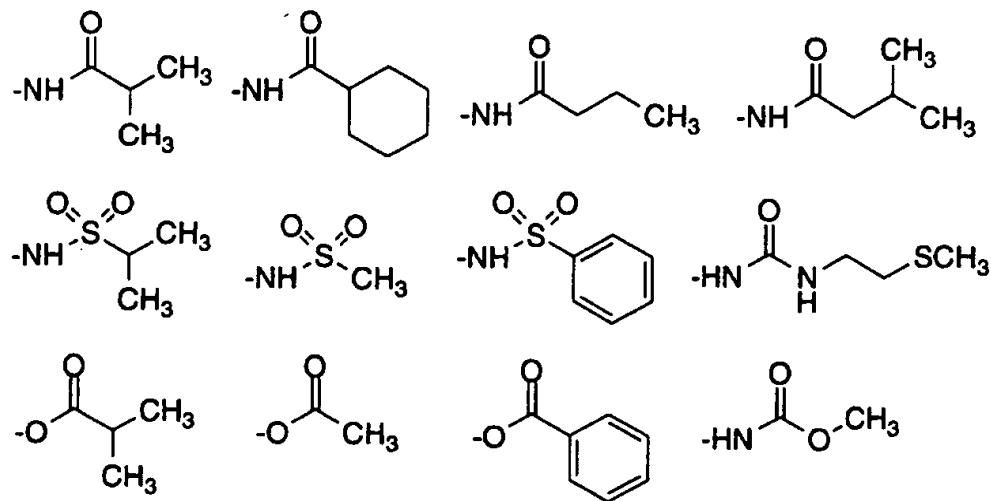
R^{4c} is selected from hydrogen, C₁-C₆ alkyl, and substituted C₁-C₆ alkyl where the substituents are selected from 1 to 5 halo, 1 to 3 hydroxy, phenyl, C₁-C₆ alkoxy carbonyl;

R^6 is selected from: hydrogen, C1-C6 alkyl, and $(CH_2)_v$ aryl, wherein the $(CH_2)_v$ and alkyl groups may be optionally substituted by $-O(R^2)$, $-S(O)_mR^2$, $-C(O)OR^2$, $-C(O)N(R^2)(R^2)$, $-SO_2N(R^2)(R^2)$, or $-N(R^2)C(O)N(R^2)(R^2)$, wherein the aryl group is selected from: phenyl, pyridyl, 1H-tetrazolyl, triazolyl, oxadiazolyl, pyrazolyl, thiadiazolyl, and benzimidazol-2-yl, which is optionally substituted with C1-C6 alkyl, C3-C6 cycloalkyl, amino, or hydroxyl;

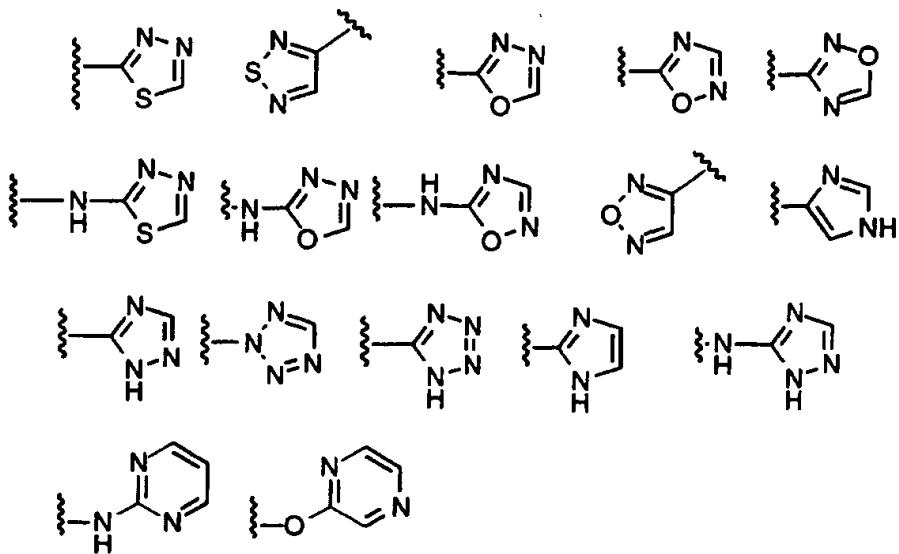
5 10 X is selected from the group consisting of: hydrogen,



- 149 -



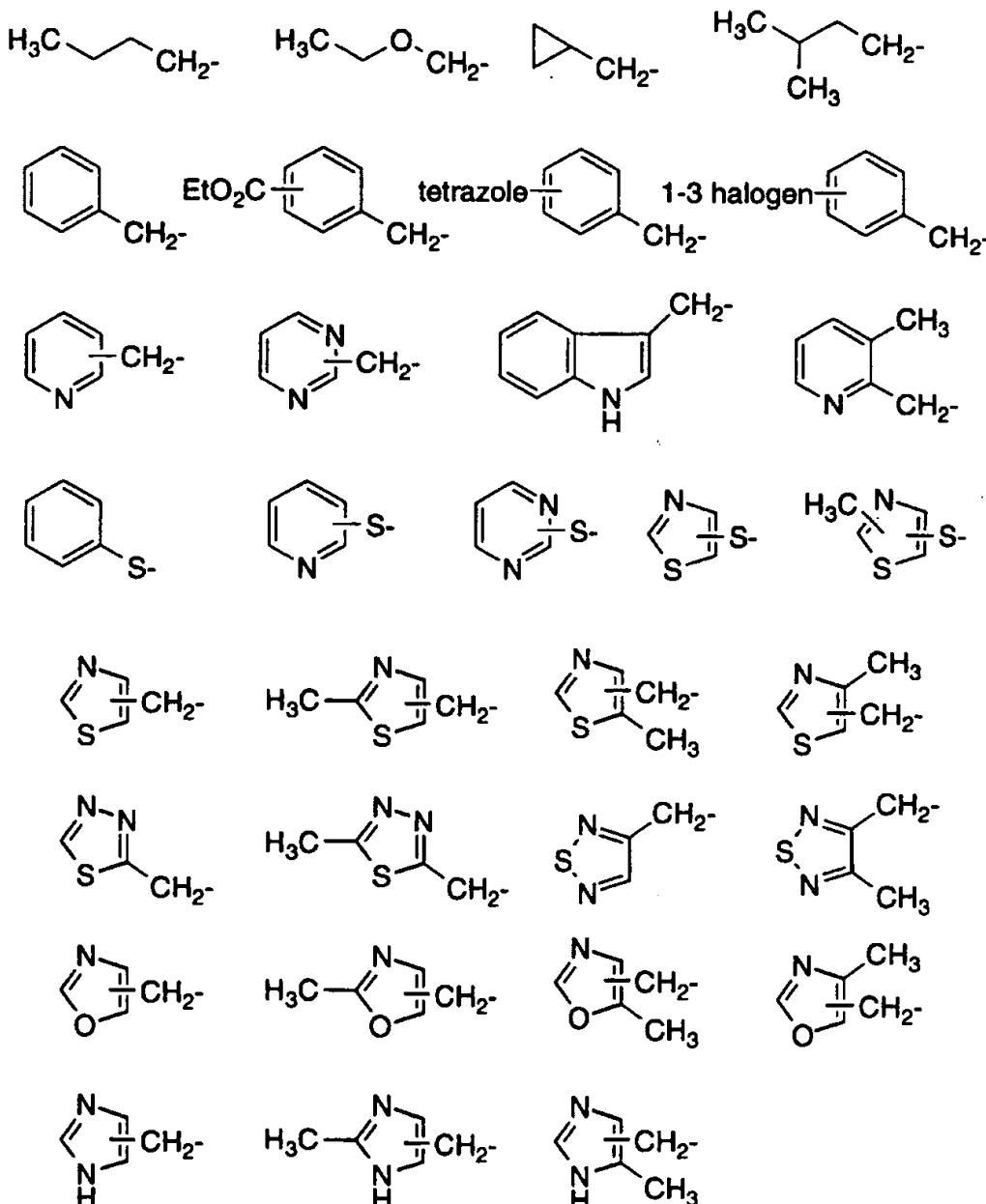
and further selected from the following group of heterocycles



5 wherein the heterocycle is optionally substituted with a substituent selected from: $-\text{N}(\text{R}^2)(\text{R}^2)$, $-\text{O}(\text{R}^2)$, C₁-C₃ alkyl, halogen, and trifluoromethyl;

- 150 -

Y is selected from the group consisting of: hydrogen,



5

or their regioisomers whereof where not specified;

- 151 -

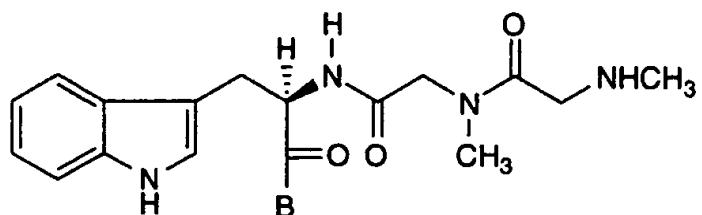
D is selected from: -N(R⁷)-, -S(O)_m-, -C(O)- and -C(H)(R⁷)-,
wherein R⁷ is selected from: -R², -(CH₂)_qaryl, -C(O)R², -SO₂R²,
-C(O)N(R²)(R²), -C(O)OR², 1-H-tetrazol-5-yl, -SO₂N(R²)aryl,
-SO₂N(R²)(R²) and the (CH₂)_q may be optionally substituted by C₁-C₄
5 alkyl, and the R² and aryl may be optionally further substituted with a
substituent selected from: -OR^{2a}, -C(O)OR^{2a}, -C(O)N(R^{2a})(R^{2a}),
halogen, -C₁-C₄ alkyl, and the aryl is selected from of triazolyl,
oxadiazolyl, 1H-tetrazolyl, and thiadiazolyl;

10 1 is 0, 1 or 2;
m is 0, 1, or 2;
q is 0, 1, 2, 3, or 4;
r is 0, 1, 2, or 3;
t is 0, 1, 2, or 3;
15 v is 0, 1, or 2;

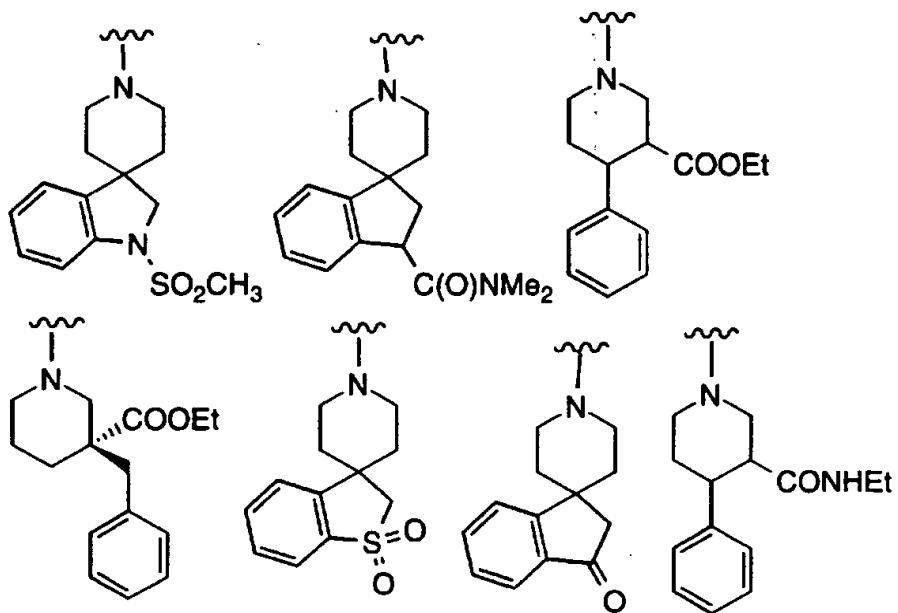
and pharmaceutically acceptable salts and individual diastereomers
thereof.

- 152 -

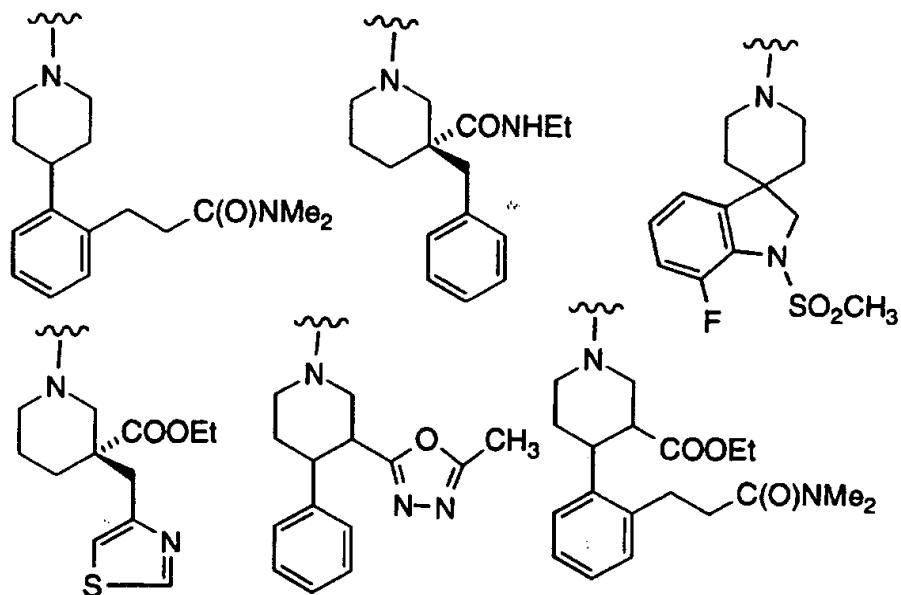
4. The compound of Claim 1 of the formula:



5 wherein B is selected from the group consisting of:



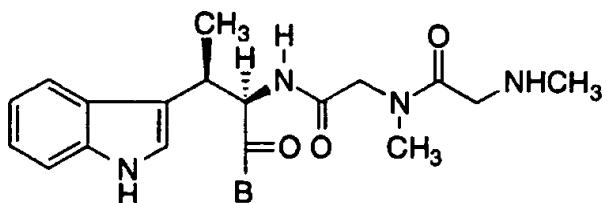
- 153 -



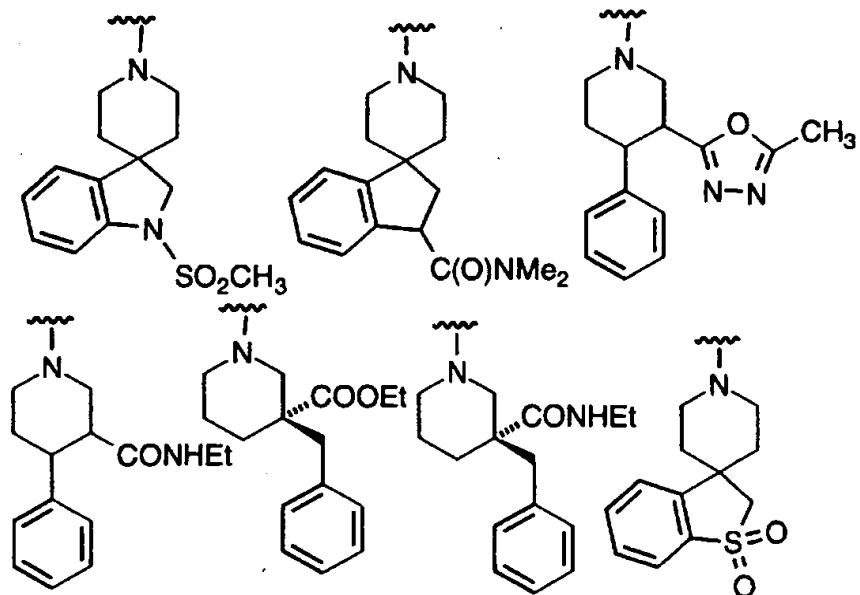
and pharmaceutically acceptable salts and individual diastereomers thereof.

- 154 -

5. The compound of Claim 1 of the formula:



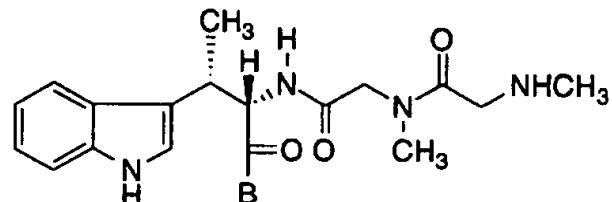
5 wherein B is selected from the group consisting of:



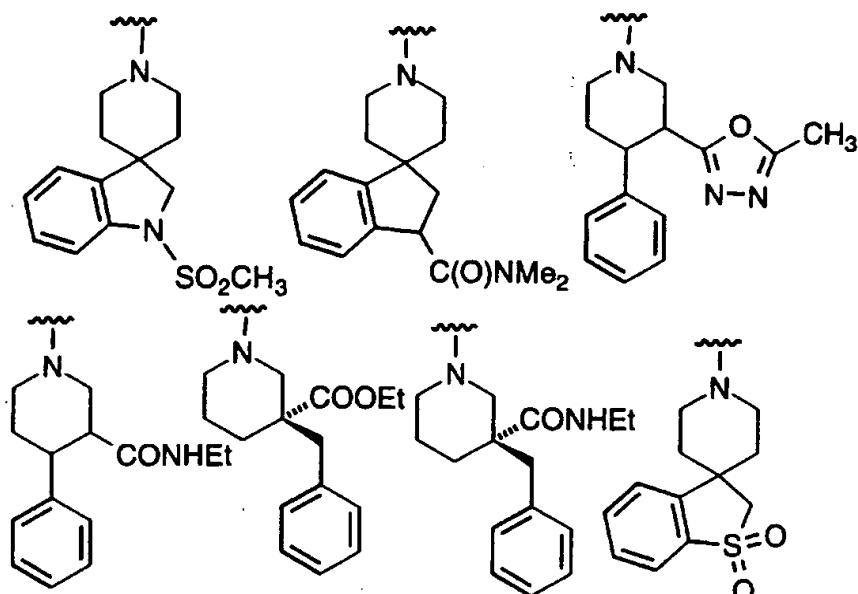
and pharmaceutically acceptable salts and individual diastereomers thereof.

- 155 -

6. The compound of Claim 1 of the formula:



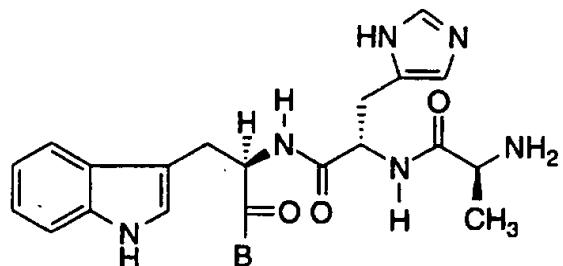
wherein B is selected from the group consisting of:



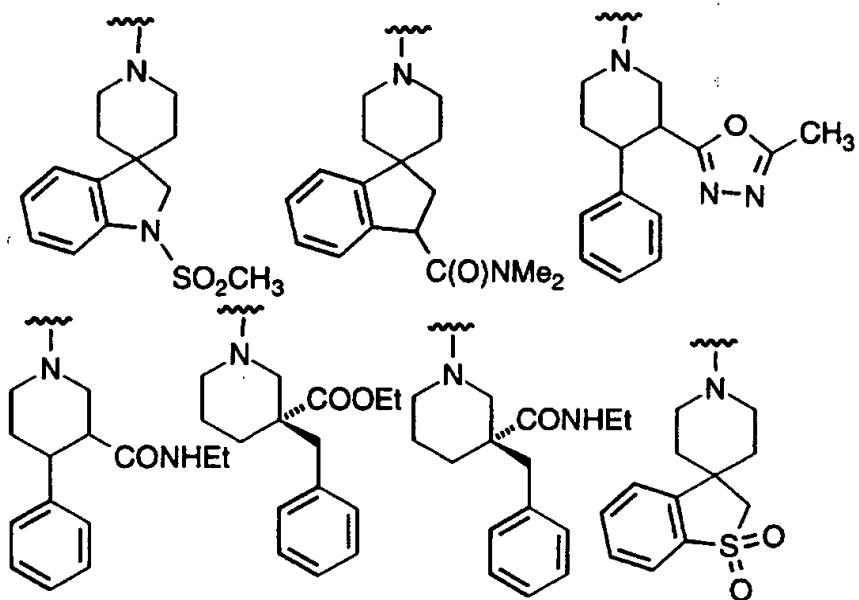
and pharmaceutically acceptable salts and individual diastereomers thereof.

- 156 -

7. The compound of Claim 1 of the formula:



wherein B is selected from the group consisting of:

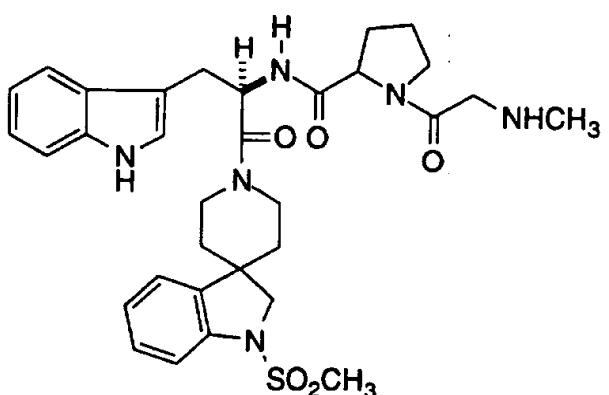
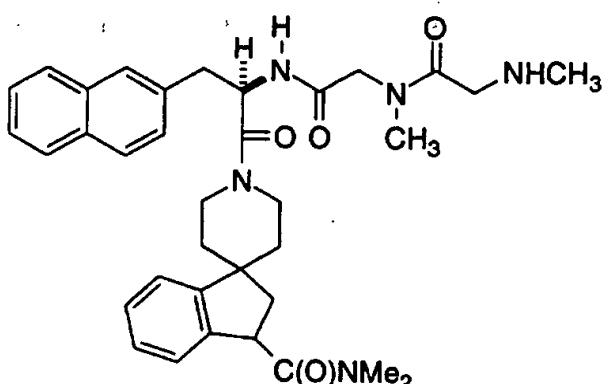
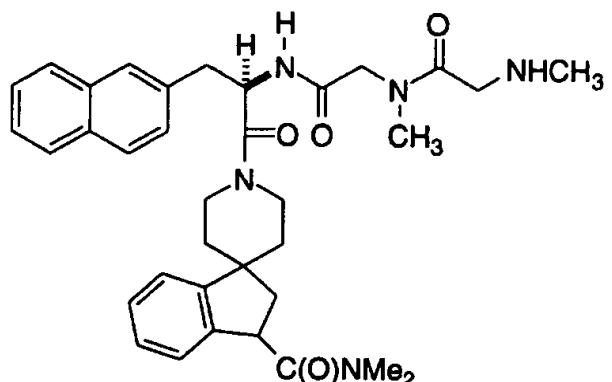


5

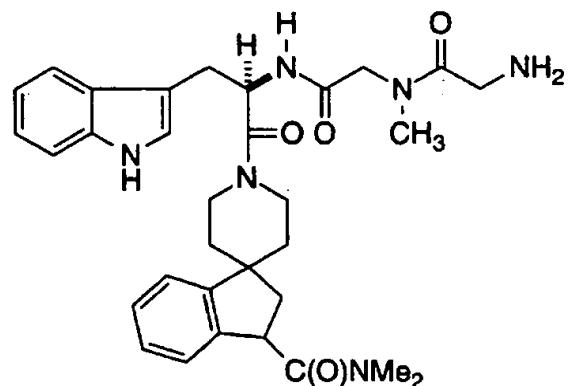
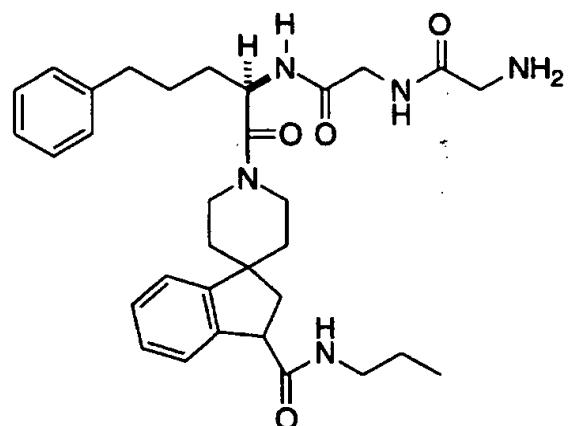
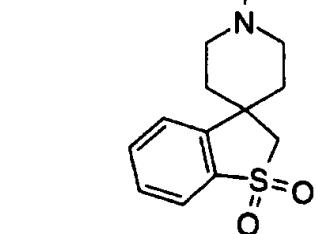
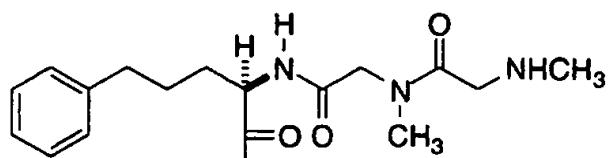
and pharmaceutically acceptable salts and individual diastereomers thereof.

- 157 -

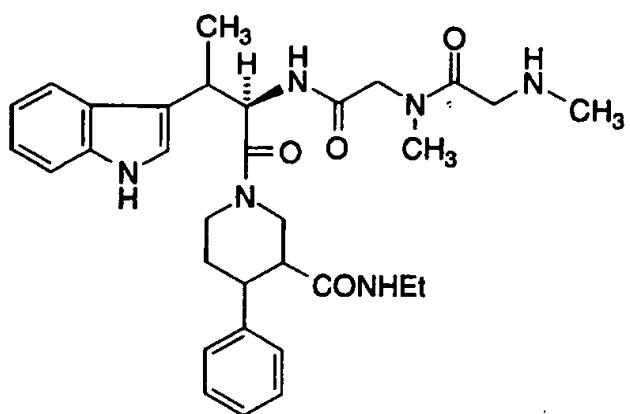
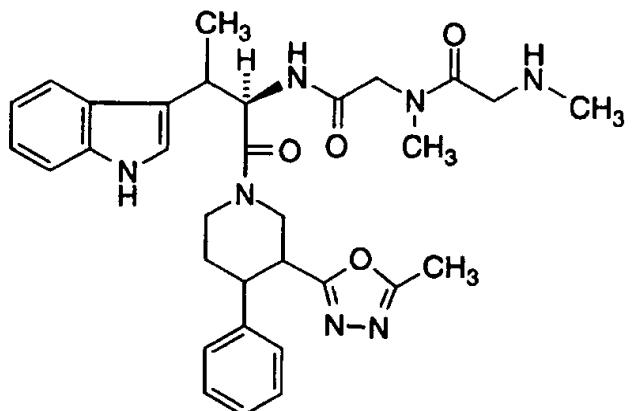
8. A compound which is selected from the group consisting of:



- 158 -



- 159 -



and pharmaceutically acceptable salts and individual diastereomers
5 thereof where not otherwise specified.

- 160 -

9. A pharmaceutical composition which comprises an inert carrier and the compound of Claim 1.

10 5 A pharmaceutical composition useful for increasing the endogenous production or release of growth hormone in a human or an animal which comprises an inert carrier and an effective amount of the compound of Claim 1 in combination with an additional growth hormone secretagogue.

10 11. A pharmaceutical composition useful for the treatment of osteoporosis which comprises a combination of a bisphosphonate compound and the compound of Claim 1.

15 12. The pharmaceutical composition of Claim 11 wherein the bisphosphonate compound is alendronate.

13. A method for increasing levels of endogenous growth hormone in a human or an animal which comprises administering to such human or animal an effective amount of the compound of Claim 1.

20 14. A method for increasing levels of endogenous growth hormone in a human or an animal which comprises administering to such human or animal an effective amount of the compound of Claim 1 in combination with an additional growth hormone secretagogue.

25 15. The method of Claim 14 wherein the additional growth hormone secretagogue is selected from the group consisting of: growth hormone releasing factor; an analog of growth hormone releasing factor; IGF-1; and IGF-2.

30 16. A method for increasing feed efficiency, promoting growth, increasing milk production and improving the carcass quality of livestock which comprises administering to such livestock an effective amount of the compound of Claim 1.

- 161 -

17. A method for the treatment of a disease or a condition which is benefited by the anabolic effects of enhanced growth hormone levels that comprises administering to a patient in need thereof 5 an effective amount the compound of Claim 1.

18. The method of Claim 17 wherein the disease or condition is selected from the group consisting of: osteoporosis; catabolic illness; immune deficiency, including that in individuals with a 10 depressed T4/T8 cell ratio; bone fracture; musculoskeletal impairment in the elderly; growth hormone deficiency in adults or in children; short stature in children; obesity; sleep disorders; cachexia and protein loss due to chronic illness such as AIDS or cancer; and the treatment of patients recovering from major surgery, wounds or burns.

15 19. A method for the treatment of osteoporosis which comprises administering to a patient with osteoporosis a combination of a bisphosphonate compound and the compound of Claim 1.

20 20. The method of Claim 19 wherein the bisphosphonate compound is alendronate.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US97/05378

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C07D 221/20; 401/12, 401/14; A61K 31/40, 31/445
US CL :546/17, 201, 210; 514/278, 323, 326

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 546/17, 201, 210; 514/278, 323, 326

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
NONEElectronic data base consulted during the international search (name of data base and, where practicable, search terms used)
CAS COMPUTER SEARCH (1966-TO DATE)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,283,241 A (BOCHIS et al) 01 February 1994, see entire document.	1-20
A	US 5,438,136 A (DEVITA et al) 01 August 1995, see entire document.	1-20
A	US 5,494,919 A (MORRIELLO et al) 27 February 1996, see entire document.	1-20

Further documents are listed in the continuation of Box C.

See patent family annex.

•	Special categories of cited documents:	
"A"	document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E"	earlier document published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L"	document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O"	document referring to an oral disclosure, use, exhibition or other means	"A" document member of the same patent family
"P"	document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search
11 JULY 1997

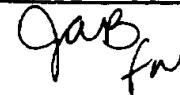
Date of mailing of the international search report

13 SEP 1997

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Authorized officer

ZINNA N. DAVIS



Facsimile No. (703) 305-3230

Telephone No. (703) 308-1235